

INDICATIONS: PROHIBIT (levamisole hydrochloride) is a broad-spectrum anthelmintic and is effective against the following nematode infections in cattle and sheep:

SHEEP:

STOMACH WORMS: *Haemonchus contortus*, *Trichostrongylus axei*, *Teladorsagia circumcincta*.
INTESTINAL WORMS: *Trichostrongylus colubriformis*, *Cooperia curticei*, *Nematodirus spathiger*, *Bunostomum trigonocephalum*, *Oesophagostomum columbianum*, *Chabertia ovina*.
LUNGWORMS: *Dictyoacaulus filaria*.

CATTLE:

STOMACH WORMS: *Haemonchus placei*, *Trichostrongylus axei*, *Ostertagia ostertagi*.
INTESTINAL WORMS: *Trichostrongylus longispicularis*, *Cooperia oncophora*, *Cooperia punctata*, *Nematodirus spathiger*, *Bunostomum phlebotomum*, *Oesophagostomum radiatum*.
LUNGWORMS: *Dictyoacaulus viviparus*.

RESIDUE WARNING: Do not administer to cattle within 48 hours of slaughter for food. Do not administer to sheep within 72 hours of slaughter for food. To prevent residues in milk, do not administer to dairy animals of breeding age.

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/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 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.

CAUTION: Muzzle foam may be observed. However, this reaction will disappear within a few hours. If this condition persists, a veterinarian should be consulted. Follow recommended dosage carefully. Consult veterinarian before using in severely debilitated animals.

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

Store between 20°-25°C (68°-77°F).

HOW SUPPLIED: 52 g (1.8 oz) and 605 g (21.34 oz)

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Manufactured for
Huvepharma, Inc.
Peachtree City, GA 30269

LOT NO.:

EXP. DATE:



TAKE TIME
OBSERVE LABEL
DIRECTIONS
M# L-6232-60
Rev. 03-2022

NDC 23243-2320-6

PROHIBIT[®]

(levamisole hydrochloride)

Soluble Drench Powder Anthelmintic

CATTLE AND SHEEP DEWORMER FOR ORAL USE

This bottle contains 544.5 grams of
levamisole hydrochloride activity.

Not For Use in Humans
Keep Out of Reach of Children
Restricted Drug (California) -
Use Only as Directed

Approved by FDA under ANADA # 200-225

Net Wt: 21.34 oz (1.3 lb.) (605 g)



DOSAGE AND ADMINISTRATION: When you are ready to deworm your cattle or sheep, add water to the powder in this bottle up to the 3 liter mark. Swirl to mix thoroughly before using. If any solution is left over, it may be stored for up to 3 months in this tightly capped bottle, shake well before using.

DATE WATER WAS ADDED TO THIS BOTTLE

Month Day Year

Administer as a single drench dose as follows:

CATTLE--2 mL per 100 lb. body weight

Weight	Drench Dosage	Bottle Will Treat
100 lb.	2 mL	1,500 head
300 lb.	6 mL	500 head
500 lb.	10 mL	300 head
700 lb.	14 mL	214 head
1,000 lb.	20 mL	150 head

SHEEP--1 mL per 50 lb. body weight

Weight	Drench Dosage	Bottle Will Treat
50 lb.	1 mL	3,000 head
100 lb.	2 mL	1,500 head
150 lb.	3 mL	1,000 head
200 lb.	4 mL	750 head

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.

NOTE: Careful weight estimates are essential for proper performance of this product. Cattle and sheep maintained under conditions of constant helminth exposure may require retreatment within two to four weeks after the first treatment.

To report suspected adverse drug events, for technical assistance or to obtain a copy of the Safety Data Sheet (SDS), contact Huvepharma, Inc. at 1-877-994-4883 or www.huvepharma.us. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or <http://www.fda.gov/reportanimalae>.



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