

PROHIBIT SOLUBLE DRENCH POWDER- levamisole hydrochloride powder, for solution

Huvepharma, Inc

PROHIBIT®
(levamisole hydrochloride)
Soluble Drench Powder

NDC 23243-2320-5

PROHIBIT®
(levamisole hydrochloride)
Soluble Drench Powder

Anthelmintic

Each packet contains 46.8 grams of levamisole hydrochloride activity.

CATTLE AND SHEEP DEWORMER FOR ORAL USE

Administer as a standard drench with standard drench syringe or administer as a concentrated drench solution with an automatic drenching syringe.

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: *Dictyocaulus filaria*.

CATTLE:

STOMACH WORMS: *Haemonchus placei*, *Trichostrongylus axei*, *Ostertagia ostertagi*.

INTESTINAL WORMS: *Trichostrongylus longispicularis*, *Cooperia oncophora*, *Cooperia punctata*,

Nematodirus spathiger, *Bunostomum phiebotomum*, *Oesophagostomum radiatum*.

LUNGWORMS: *Dictyocaulus viviparus*.

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 WEIGHT: 1.8 oz (52 g)

HUVEPHARMA®

Manufactured for Huvepharma, Inc.

Peachtree City, GA 30269

®Registered Trademark of Huvepharma, Inc.

DOSAGE AND ADMINISTRATION

CATTLE-STANDARD DRENCH

SOLUTION: Place the contents of this packet in a 1 quart (32 fl. oz.) container, fill with water, swirl until dissolved. Administer as a single drench dose according to the following table:

Weight	Drench dosage	Packet Will Treat
200 lb.	1/2 fl. oz.	64 head
400 lb.	1 fl. oz.	32 head
650 lb.	1 1/2 fl. oz.	21 head
800 lb.	2 fl. oz.	16 head

SHEEP-STANDARD DRENCH SOLUTION:

Place the contents of this packet in a 1 gallon (128 fl. oz.) container, fill with water, swirl until dissolved. Administer as a single drench dose according to the following table:

Weight	Drench Dosage	Packet Will Treat
50 lb.	1/2 fl. oz.	256 head
100 lb.	1 fl. oz.	128 head
150 lb.	1 1/2 fl. oz.	84 head
200 lb.	2 fl. oz.	64 head

CONCENTRATED DRENCH SOLUTION: For use with automatic syringe. Place the contents of this packet in a standard household measuring container and add water to the 8 3/4 fl. oz. level; or use the measuring container available from your supplier and add water to the mark. Swirl until dissolved. Give 2 ml (milliliter) per 100 lb. body weight. Refer to the table above for the number of cattle this packet will treat.

CONCENTRATED DRENCH SOLUTION: For use with automatic syringe. Place the contents of this packet in a standard household measuring container and add water to the 17 1/2 fl. oz. level. Swirl until dissolved. Give 2 ml per 50 lb. body weight. Refer to the table above for the number of sheep this packet will treat.

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. Prepare solutions as needed. However, excess solutions may be stored in clean closed containers up to 90 days without loss of anthelmintic activity. Cattle and Sheep maintained under conditions of constant helminth exposure may require re-treatment within two to four weeks after the first treatment.

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.

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>.

Store between 20–25°C (68–77°F). MF: L-6232-05 Rev: 03-2022

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

LOT

EXP.

TAKE TIME



OBSERVE LABEL
DIRECTIONS

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)

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 circumscincta*.

INTESTINAL WORMS: *Trichostrongylus colubriformis*, *Cooperia curticei*, *Nematodirus spathiger*,

Bunostomum trigonocephalum, *Oesophagostomum columbianum*, *Chabertia ovina*.

LUNGWORMS: *Dictyocaulus filaria*.

CATTLE:

STOMACH WORMS: *Haemonchus placei*, *Trichostrongylus axei*, *Ostertagia ostertagi*.

INTESTINAL WORMS: *Trichostrongylus longispicularis*, *Cooperia oncophora*, *Cooperia punctata*,

Nematodirus spathiger, *Bunostomum phiebotomum*, *Oesophagostomum radiatum*.

LUNGWORMS: *Dictyocaulus 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

MF# L-6232-60

Rev. 03-2022

LOT.: **EXP. DATE**

TAKE TIME



OBSERVE LABEL
DIRECTIONS

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
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		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.

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LUNGWORMS: *Dictyoaculus filaria*.

CATTLE:

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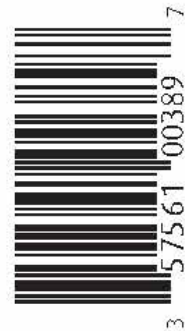
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HOW SUPPLIED: 52 g (1.8 oz) and 605 g (21.34 oz)

LOT **EXP.**

Rev: 03-2022



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANADA	ANADA200225	04/20/2022	

Labeler - Huvepharma, Inc (619153559)

Registrant - Huvepharma EOOD (552671651)

Revised: 12/2023

Huvepharma, Inc