SAFE-GUARD- fenbendazole suspension
Schering Corporation

---

safe-guard®
(fenbendazole)
Dewormer

INDICATIONS:

Goats - 2.3 mg/lb (5 mg/kg) body weight for the treatment and control of:

Stomach worms (adults): *Haemonchus contortus* and *Teladorsagia circumcincta*.

DIRECTIONS: Administer orally. The recommended dose of 2.3 mg/lb (5 mg/kg) is achieved when 2.3 mL of the drug are given for each 100 lb body weight.

Dosing Examples for Goats:

Body Weight	Amount
25 lb	0.6 mL
50 lb	1.2 mL
75 lb	1.7 mL
100 lb	2.3 mL
125 lb	2.9 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.

Fenbendazole (active ingred.) made in China. Formulated in France.

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, or http://www.fda.gov/reportanimalae.

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, 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.

Store at or below 25°C (77°F). Protect from freezing. Shake well before use.

Restricted drug (California) - Use only as directed.

Distributed by: Intervet Inc
(d/b/a Merck Animal Health)

Madison, NJ 07940 ©2020 Intervet Inc.,
a subsidiary of Merck & Co. Inc.

LOT NO:

EXP
DATE:

Rev. 09/20

Approved by FDA under
NADA # 128-620

387770 R11

PRINCIPAL DISPLAY PANEL - 125 mL Bottle Label

safe-guard®
(fenbendazole)

Dewormer
for Goats

Suspension 10%
(100mg/mL)

Withdrawal Periods and Residue Warnings: Goats must not be slaughtered for human consumption within 6 days following treatment. Because a milk discard time has not been established, do not use in lactating goats.

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

MERCK
Animal Health

125 mL (4.2 fl oz)

368607 R6

safe-guard®
(fenbendazole)

Dewormer

for Goats

Suspension 10%
(100mg/mL)

***Withdrawal Periods and Residue Warnings*:** Goats must not be slaughtered for human consumption within 6 days following treatment. Because a milk discard time has not been established, do not use in lactating goats.

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

MERCK Animal Health

368607 R6 **125 mL** (4.2 fl oz)

INDICATIONS: Goats - 2.3 mg/lb (5 mg/kg) body weight for the treatment and control of:
Stomach worms (adults): *Haemonchus contortus* and *Teladorsagia circumcincta*.

DIRECTIONS: Administer orally. The recommended dose of 2.3 mg/lb (5 mg/kg) is achieved when 2.3 mL of the drug are given for each 100 lb body weight.

0 21784 04537 7

Approved by FDA under
NADA # 128-620

387770 R11

MERCK Animal Health

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, or http://www.fda.gov/reportanimalae.

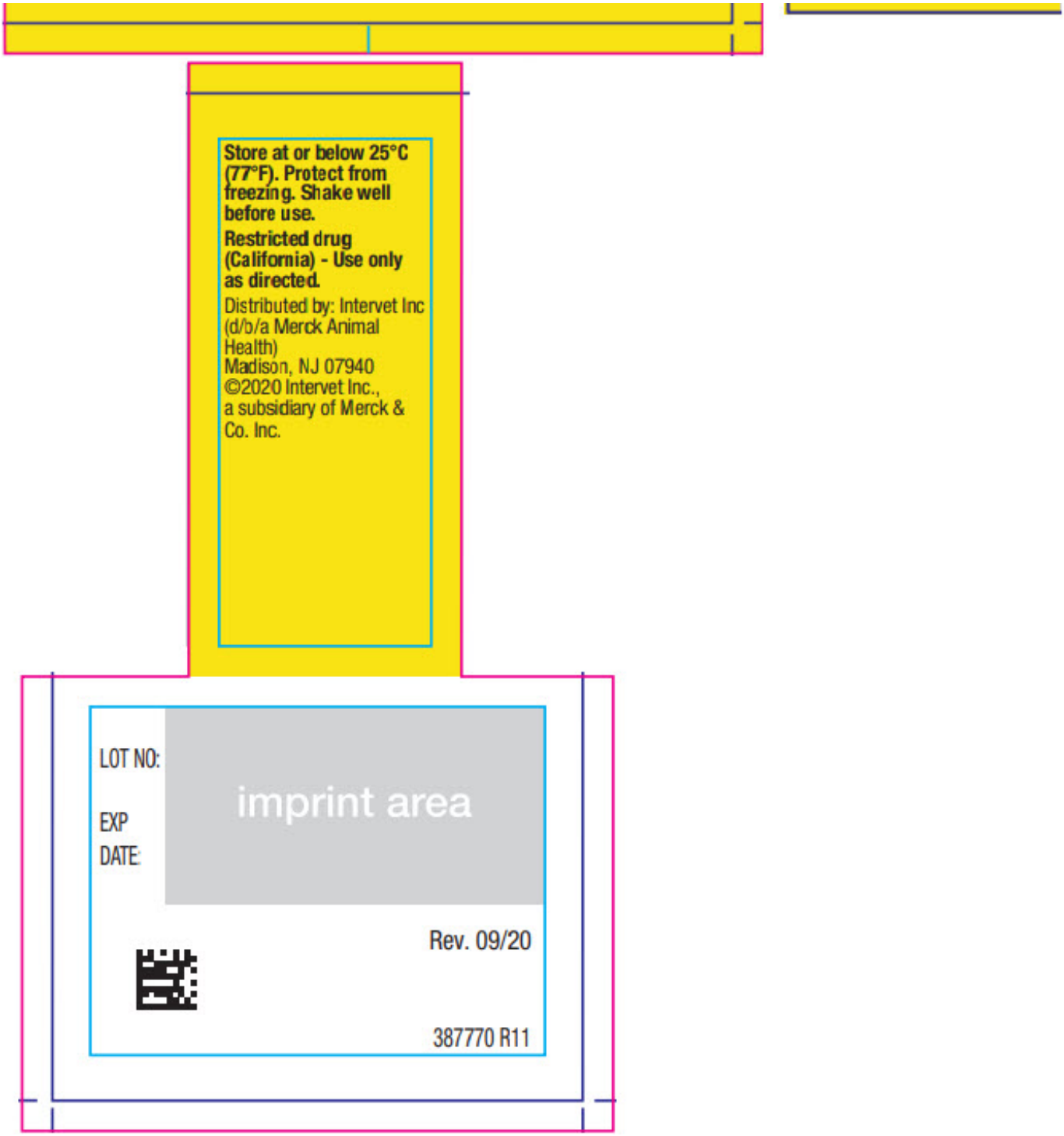
OTHER WARNINGS: Parasite resistance may develop to any dewormer, and has been reported for most classes of dewormers. Treatment with a dewormer used in conjunction with parasite management practices appropriate to the geographic area and the animal(s) to be treated may slow the development of parasite resistance. Fecal examinations or other diagnostic tests and parasite management history should be used to determine if the product is appropriate for the herd, 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.

Dosing Examples for Goats:

Body Weight	Amount
25 lb	0.6 mL
50 lb	1.2 mL
75 lb	1.7 mL
100 lb	2.3 mL
125 lb	2.9 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.

Fenbendazole (active ingred.) made in China. Formulated in France.



SAFE-GUARD

fenbendazole suspension

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:57926-089
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:57926-089-01	125 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA128620	09/20/1983	

Labeler - Schering Corporation (001317601)

Establishment

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

Establishment

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

Revised: 9/2021 Schering Corporation