SAFE-GUARD- fenbendazole suspension
Merck Sharp & Dohme Corp.

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safe-guard®
(fenbendazole)
Dewormer

INDICATIONS:

Beef and Dairy Cattle - 2.3 mg/lb (5 mg/kg) body weight 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*).

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) of body weight is achieved when 2.3 mL of the drug are given for each 100 lb body weight.

EXAMPLES:

Cattle:		**Goats:**	
Body Weight	**Amount**	**Body Weight**	**Amount**
100 lb	2.3 mL	25 lb	0.6 mL
200 lb	4.6 mL	50 lb	1.2 mL
300 lb	6.9 mL	75 lb	1.7 mL
400 lb	9.2 mL	100 lb	2.3 mL
500 lb	11.5 mL	125 lb	2.9 mL
1000 lb	23.0 mL		
1500 lb	34.5 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, 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.

Fenbendazole (active ingred.) made in: see imprint.
Formulated in France.
Distributed by: Intervet Inc.
(d/b/a Merck Animal Health), Rahway, NJ 07065

Restricted drug (California) - Use only as directed.

Approved by FDA under NADA # 128-620

Rev. 04/2024
217157 R1

PRINCIPAL DISPLAY PANEL - 3,785 mL Bottle Label

safe-guard®
(fenbendazole)

Dewormer

for Beef & Dairy Cattle
and Goats

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. Goats must not be slaughtered for human consumption

within 6 days following last treatment with this drug product. 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.

1 Gallon (3,785 mL)

MERCK
Animal Health

LOT NUMBER:
EXPIRATION DATE:

386508 R4

safe-guard® (fenbendazole)

Dewormer

for Beef & Dairy Cattle and Goats

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.

Goats must not be slaughtered for human consumption within 6 days following last treatment with this drug product. 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.

LOT NUMBER:

imprint area

EXPIRATION DATE:

1 Gallon (3,785 mL)

MERCK Animal Health

386508 R4

INDICATIONS: Beef and Dairy Cattle - 2.3 mg/lb (5 mg/kg) body weight 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*).
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) of body weight is achieved when 2.3 mL of the drug are given for each 100 lb body weight.

EXAMPLES:

Cattle:		Goats:	
Body Weight	Amount	Body Weight	Amount
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1000 lb	23.0 mL		
1500 lb	34.5 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, 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.
Fenbendazole (active ingred.) made in China. Formulated in France.
Distributed by: Intervet Inc (d/b/a Merck Animal Health), Madison, NJ 07940
Approved by FDA under NADA # 128-620

Rev. 09/20

0 21784 46710 0

MERCK Animal Health

367235 R3

PRINCIPAL DISPLAY PANEL - 1,000 mL Bottle Label

safe-guard®
(fenbendazole)

Dewormer

for Beef & Dairy Cattle
and Goats

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. Goats must not be slaughtered for human consumption within 6 days following last treatment with this drug product. 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.

1,000 mL (33.8 fl oz)

MERCK
Animal Health

361394 R9

safe-guard®
(fenbendazole)

Dewormer

for Beef & Dairy Cattle and Goats

Suspension 10% (100 mg/mL)

Withdrawal Periods and Residue Warnings:
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Consult your veterinarian for assistance in the diagnosis, treatment and control of parasitism.

1,000 mL (33.8 fl oz)

MERCK Animal Health

361394 R9

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EXAMPLES:

Cattle:		Goats:	
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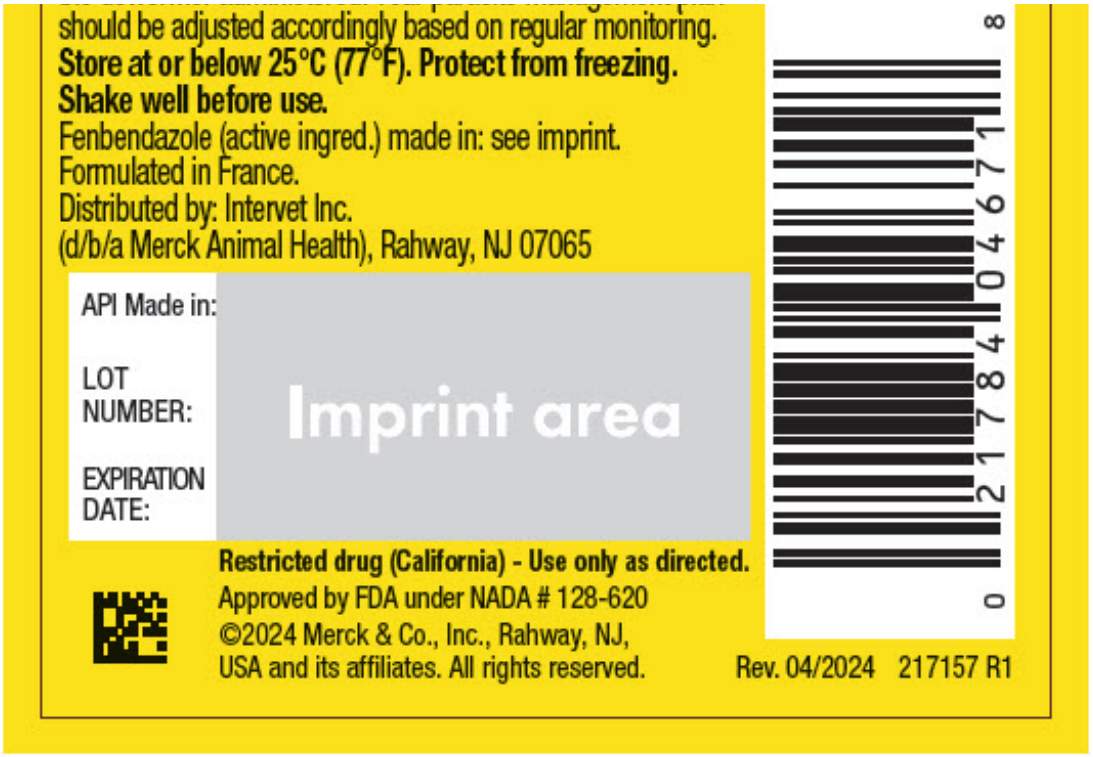
MERCK Animal Health



SAFE-GUARD

fenbendazole suspension

Product Information

Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:57926-088
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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57926-088-01	3785 mL in 1 BOTTLE		
2	NDC:57926-088-02	1000 mL in 1 BOTTLE		
3	NDC:57926-088-03	10000 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 - Merck Sharp & Dohme Corp. (001317601)

Revised: 11/2024 Merck Sharp & Dohme Corp.