

PRO-TECT HOT SPOT SKIN MEDICATION- benzalkonium chloride liquid
Straight Arrow Products, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredient:

Benzalkonium chloride

Purpose:

Purpose: Pro-Tect Hot Spot Medication for dogs is formulated to be effective for the treatment of skin inflammations. Specifically kills staphylococcus intermedius; bacteria most commonly associated with "Hot Spots" on dogs. Also, treats dogs for the secondary skin infections caused by viral, fungal or bacterial organisms. **Pro-Tect Hot Spot** is an effective first aid treatment for superficial cuts, scrapes, and abrasions. Best of all, **Pro-Tect Hot Spot** contains no alcohol and is a non-stinging formula.

Symptoms of Hot Spots: Hot Spots are usually accompanied by incessant rubbing, scratching, and biting due to intense itching. Other signs are moist reddened areas, hair loss, and flaky skin. Hot spots are usually warm to the touch.

Directions for Use:

Trim the hair around the sore. Wash the area with mild soap and water, rinse and pat dry. Apply Pro-Tect Hot Spot to affected area. Does not require dabbing on with a cotton ball. Do not use on puppies under 6 weeks of age.

General Precautions:

Consult a veterinarian if any problems persist or are of concern.

Caution: KEEP OUT OF REACH OF CHILDREN



Pro-Tect® Hot Spot Skin Medication for Dogs

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Active Ingredient: Benzalkonium Chloride

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UPC Code

Made and Distributed
in the U.S.A. by:
Straight Arrow Products, Inc.
P.O. Box 20350
Lehigh Valley, PA 18002-0350
Galloping Horses © 1992 Straight Arrow Products, Inc.

4 Fl. Oz. (118 ml)



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benzalkonium chloride liquid

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:62001-0317
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.001 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	

IMIDUREA (UNII: M629807ATL)
DIDECYLDIMONIUM CHLORIDE (UNII: JXN4009Y9B)
OCTOXYNOL 9 (UNII: 7JPC6Y25QS)
QUATERNIUM-15 (UNII: E40U03LEM0)
METHYLPARABEN (UNII: A2I8C7HI9T)
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)
ALLANTOIN (UNII: 344S277G0Z)
PROPYLPARABEN (UNII: Z8IX2SC1OH)
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)
TROLAMINE (UNII: 9O3K93S3TK)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62001-0317-1	118 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/25/2003	

Labeler - Straight Arrow Products, Inc. (061580593)

Registrant - Straight Arrow Products, Inc. (061580593)

Establishment

Name	Address	ID/FEI	Business Operations
Straight Arrow Products, Inc.		061580593	manufacture, pack

Establishment

Name	Address	ID/FEI	Business Operations
Lonza Biologics, Inc.		093149750	api manufacture

Revised: 7/2025

Straight Arrow Products, Inc.