## HARTZ GROOMERS BEST- hydrocortisone spray The Hartz Mountain Corporation

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.

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## **Drug Facts**

#### **Active Ingredient**

0.5% Hydrocortisone, USP

#### **Purpose**

Anti-itch

**Uses** to soothe • Itching •Flea bites •Dermatitis •Skin Irratations •Eczema

## Warnings

#### For external use on adult dogs and cats only.

Keep out of the reach of children and animnals to avoid unintended consumption.

**Do not use** •In eyes or nose •not for prolonged use •do not apply to large areas of the body •do not use where infection (pus) is present, since the drug may allow the infection to be spread •on pregnant animals • if product appears to be tasmpered with

stop use and ask a veterinarian if redness, irritations, or swelling persists or increases

**Directions:** Shake well before use. Hold sprayer 4 to 6 inches from affected area and apply directly. Apply 2-3 times daily to irritated area until relief is achieved. For best results, cleanse affected area prior to use when using this product. Avoid contact with eyes

*Other information*Store product in cool, dry place. Avoid temperatures above 86F

#### **Inactive Ingredients**

Aloe Vera Gel, Deionized Water, Disodium EDTA, Propylene Glycol



#### HARTZ GROOMERS BEST

hydrocortisone spray

Product Information						
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:43201-3056			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety						
Ingredient Name	Basis of Strength	Strength				
HYDROCORTISONE (UNII: WI4X0 X7BPJ) (HYDROCORTISONE - UNII:WI4X0 X7BPJ)	HYDROCORTISONE	0.5 g in 100 mL				

Inactive Ingredients				
Ingredient Name	Strength			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
WATER (UNII: 059QF0KO0R)				
EDETATE DISO DIUM (UNII: 7FLD91C86K)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				

Packaging								
# Item Code	Package Description	Market	ing Start Date	M	larketing End Date			
1 NDC:43201-3056-5	147 mL in 1 BOTTLE, SPRAY							
Marketing Information								
Marketing Category	Application Number or Monograph Citation		Marketing Start Date		Marketing End Date			

# $\pmb{Labeler} \textbf{ - The Hartz Mountain Corporation (058109158)}$

unapproved drug other

Revised: 4/2013 The Hartz Mountain Corporation

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