HORSEMANS DREAM FUNG-A-WAY - benzalkonium chloride solution
SUMMIT INDUSTRIES

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.

HORSEMAN'S DREAM FUNG-A-WAY TOPICAL FUNGICIDE SOLUTION

Drug Facts

Active Ingredients	Purpose
Benzalkonium chloride 0.15%	Skin wound cleaner

purpose: skin wound cleaner

Uses:

For use on horses, dogs, and cats, as an aid in the control of summer itch, girth itch and ringworm

Warnings

For external veterinary use only

Nor for human use

Not for use on animals intended for food

When using this product

do not get into eyes or mucous membranes. If contact occurs, flush immediately with water.

Stop use and consult your veterinarian if

no improvement is noted within seven days.

Keep out of reach of children. In case of contact with eyes or mucous membranes, obtain medical attention for eye inflammation.

Directions

soak affected area liberally with topical fungicide solution.

Apply daily until hair begins to grow.

Leave treated area uncovered

Rinse treated ares with clear water before reapplying

Results should be apparent in a matter of days

Note: efficiency is neutralized by soap or detergent residues.

Inactive Ingredients:

Deionized water, carbamide, allantoin, quaternium-15, FC and C yellow no. 5, FD and C red no. 40

HORSEMAN'S DREAM

FUNG-A-WAY

Topical Fungicide Solution

-Aids in control of

-Summer itch

-girth itch-

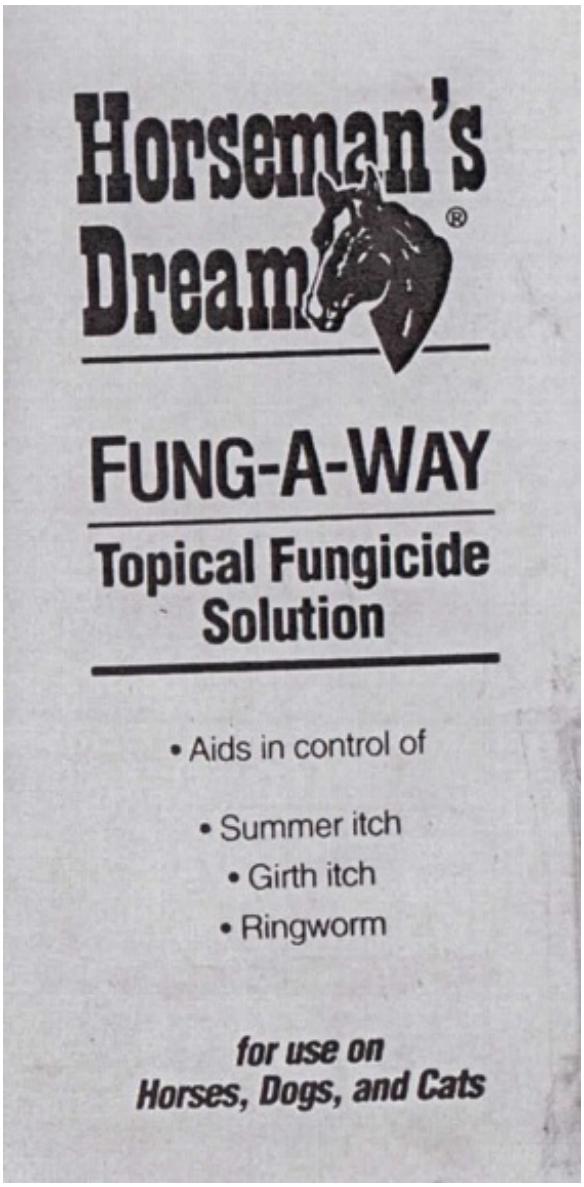
-ringworm

for use on Horses, dogs, and cats

Manufactured Exclusively For:

Horseman's Dream Inc.

Forth Worth, TX 76126



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Manufactured Exclusively For:
HORSEMAN'S DREAM INC.
Fort Worth, TX 76126

HORSEMANS DREAM FUNG-A-WAY
benzalkonium chloride solution

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:12090-1040
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.15 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
UREA (UNII: 8W8T17847W)	
ALLANTOIN (UNII: 344S277G0Z)	
QUATERNIUM-15 (UNII: E40U03LEM0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:12090-1040-8	236 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/17/2010	

Labeler - SUMMIT INDUSTRIES (003279189)

Registrant - SUMMIT INDUSTRIES (003279189)

Establishment

Name	Address	ID/FEI	Business Operations
GDMI, INC		040646846	manufacture

Revised: 9/2010 SUMMIT INDUSTRIES