FUNGUS FIGHTER - phenol liquid Continental Manufacturing Chemist, 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.

Indications:

For use on Cattle, Sheep, Horses, Dogs and other large animals as an aid in the control of ringworm, summer itch, girth itch, foot rot and other fungal infections. Do not use on cats.

Directions for use:

Shake well. Clean the area to be treated, and apply Fungus Fighter directly to the affected area, soaking completely. Let dry. Repeat application daily until condition improves, up to 4 consecutive days. For Foot Rot, apply directly to the affected area, soaking completely. Use daily until condition improves.

Precautions:

Use only as directed. For external animal use only. **KEEP OUT OF REACH OF CHILDREN.** Avoid contact with eyes and mucous membranes. If irritation develops, or if condition does not improve, discontinue use and consult veterinarian. Not intended for use on animals in commercial food production.

Warning:

Flammable!Keep away from excessive heat or open flame.

Inactive Ingredients:

alcohol (UNII: 3K9958V90M)

Ethyl Alcohol, o-Phenylphenol, Polyvinylprrolidone, Propylene Glycol, Purified Water

FUNGUS FIGHTER				
phenol liquid				
Product Information				
Product Type	OTC ANIMAL DRUG	Ite m	Code (Source)	NDC:49794-017
Route of Administration	TOPICAL			
A - 4' T 1' 4/A - 4' D.K - 1'				
Active Ingredient/Active Moiety				
Ingredie	nt Name		Basis of Strength	Strength
phenol (UNII: 339NCG44TV) (Phenol -	UNII:339 NCG44TV)		phenol	0.50 mg in 100 mL
Inactive Ingredients				
	Ingredient Name			Strength

propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	
ORTHOPHENYLPHENOL (UNII: D343Z75HT8)	
PO VIDONE K30 (UNII: U725QWY32X)	

F	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49794-017-32	946 mL in 1 BOTTLE, PLASTIC		
2	NDC:49794-017-28	3785 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/12/2010	

Labeler - Continental Manufacturing Chemist, Inc. (005278007)

Establishment				
Name	Address	ID/FEI	Business Operations	
Continental Manufacturing Chemist, Inc.		005278007	manufacture	

Revised: 3/2011 Continental Manufacturing Chemist, Inc.