FAST RESULTS GENITAL HERPES TREATMENT - benzalkonium chloride, allantoin liquid Merix Pharmaceutical Corp.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUGS FACTS

ACTIVE INGREDIENTS:

Benzalkonium chloride 0.13 % Allantoin 0.5 %

USES:

For the relief of symptoms associated with genital herpes First Aid to guard against secondary skin infection

WARNINGS:

Use only as directed. For external use only. Do not use if allergic to daisies. Stop use and ask doctor if condition worsens or last more than 10 days. **Do not use on yeast infections.**

KEEP OUT OF REACH OF CHILDREN:

Keep this and all drugs out of reach of children.

In case of accidental ingestion other then intended use, seek professional assistance or call a poison control center.

DIRECTIONS:

Adults or adolescents 12 or older apply to clean dry area free of soap or cleanser residue.

Apply to area as need 3-4 times daily. Do no use cotton to apply. Allow to dry.

Other Ingredients:

Purified water, Viracea (a patented blend of Bensalkonium chloride and Echinacea) Methylparaben, Propylparaben, Potassium sorbate.

Fast Results label



FAST RESULTS GENITAL HERPES TREATMENT

benzalkonium chloride, allantoin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63287-425
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Benzalkonium Chloride (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	1.3 mL in 1 mL		
Allantoin (UNII: 344S277G0Z) (Allantoin - UNII:344S277G0Z)	Alla nto in	5 mL in 1 mL		

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Methylparaben (UNII: A2I8C7HI9T)			
Potassium Sorbate (UNII: 1VPU26JZZ4)			
Propylparaben (UNII: Z8IX2SC1OH)			
Echinacea purpurea flowering top (UNII: 2EMS3QFX65)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63287-425-25	1 in 1 BOX		
1		25 mL in 1 BOTTLE, SPRAY		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/01/2006		

Labeler - Merix Pharmaceutical Corp. (158385687)

Registrant - Topical Pharmaceutiocals Inc. (831530683)

Establishment					
Name	Address	ID/FEI	Business Operations		
Topical Pharmaceutiocals Inc.		831530683	manufacture		

Revised: 5/2012 Merix Pharmaceutical Corp.