RELEEV COLD SORE TREATMENT- benzalkonium chloride 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.

RELEEV 1 day Cold sore Symptom Treatment

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Cold Sore/ Fever Blister Treatment

Uses:

Treats cold sores/ fever blisters. For the pain, tingling, itching, & burning associated with cold sores. May be used as an antiseptic to help cleanse or dry cold sores & fever blisters. May be used inside the mouth on sores.

Warnings:

For external use only

Do Not Use:

If you have ever had an allergic reaction to this product or any of its ingredients.
Do not use for yeast infections (may be used in conjunction with yeast medication).
Avoid contact with eyes.
If condition worsens or does not improve, contact a health care professional.

When Using This Product

• Use only as directed • Brief tingling may occur.

Keep this and all drugs out of reach of children. In case of accidental ingestion other than intended use, seek professional assistance or call a poison control center.

Directions:

SHAKE WELL (Adults and children 12 years of age or older) apply to clean, dry, affected area 3-4 times per day by dabbing and pressing solution into the area well. Allow to dry. Best when used at first sign of outbreak. Do not use cotton applicator.

Inactive Ingredients:

Echinacea Purpurea [Viracea®, proprietary blend of Benzalkonium Chloride and

Echinacea herb], Methyl Cellulose, Methyl Paraben, Potassium Sorbate, Propyl Paraben, Purified Water,

Other Information:

We suggest that you not consume alcohol during an outbreak or when using this product. Due to the natural ingredients in this product, colour may vary. Product dries clear.

Packaging



Other Information: We suggest that you not consume alcohol during an outbreak or when using



RELEEV COLD SORE TREATMENT benzalkonium chloride liquid **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:63287-419 **Route of Administration** TOPICAL **Active Ingredient/Active Moiety Ingredient Name** Basis of Strength Strength BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -**BENZALKONIUM** 1.3 mg UNII:7N6JUD5X6Y) CHLORIDE in 1 mL **Inactive Ingredients** Strength **Ingredient Name** HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35)

ECHINACEA PURPUREA FLOWERING TOP (UNII: 2EMS3QFX65)

METHYLPARABEN (UNII: A2I8C7HI9T)				
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)				
WATER (UNII: 059QF0KO0R)				
PROPYLPARABEN (UNII: Z8IX2SC10H)				

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:63287- 419-01	1 in 1 BLISTER PACK	06/01/2006			
1		1 in 1 POUCH				
1		1 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product				
2	NDC:63287- 419-03	1 in 1 BLISTER PACK	06/01/2006	08/31/2012		
2		3 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product				
3	NDC:63287- 419-06	1 in 1 BLISTER PACK	06/01/2006			
3		6 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product				
4	NDC:63287- 419-33	1 in 1 BLISTER PACK	08/21/2018			
4		3 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product				
Marketing Information						
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
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Labeler - Merix Pharmaceutical Corp. (158385687)

Revised: 7/2022

Merix Pharmaceutical Corp.