

SHING-RELEEV - 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.

DRUGS FACTS

ACTIVE INGREDIENTS:

Benzalkonium Chloride .13%

Allantoin .5%

Benzyl Alcohol .5%

USES:

For the relief of symptoms associated with shingles including pain, Burning, Itching and tingling First aid to help guard against secondary skin infection due to shingles.

WARNINGS:

For external use only. Not for ingestion.

Do not use- in yeast infections- do not spray directly on the eyes

When using this product, may tingle on contact.

Stop used and ask doctor if - condition worsens- symptoms

last more than 7 days .

KEEP OUT OF REACH OF CHILDREN:

If swallowed get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Use at first sign of irritation or itching.

Adults and children 12 years or older. Clean without soap.
apply liberally to clean dry area free of soap or cleanser residue.

Apply to area as needed 3-4 times daily.

Do not use cotton applicator.

May be used with sterile bandage after area is dry.

Other Ingredients

.Methylparaben Potassium Sorbate (natural preservative).

Propylparaben , Viracea (proprietary Echinacea purpurea extract),
water (purified)

Shing carton



Drug Facts

Active ingredient	Purpose
benzalkonium chloride 0.13%	topical antiseptic
allantoin	skin protectant
benzyl alcohol	external analgesic

Uses

- For the relief of symptoms associated with shingles.
- First aid to help guard against secondary skin infection due to shingles.

Warnings

For external use only. Not for ingestion.

Do not use in yeast caused infections • do not spray directly on the eye.

When using this product, may sting on contact.

Stop use and ask doctor if condition worsens; symptoms last more than 7 days.

Keep out of the reach of children. If swallowed, get medical help or call a Poison Control Center right away.

Directions

- Use at first sign of irritation or itching.
- Adults or children 2 years of age or older: Apply liberally to clean dry area free of any soap or cleanser residue. Apply to area as needed 3-4 times daily. Do not use cotton applicator. May be covered with loose sterile bandage after area is dry.

Inactive ingredients

Methyl Paraben (natural preservative from blueberries), Potassium Sorbate (natural preservative), Propyl Paraben (natural preservative from blueberries), Vinaces® (proprietary Echinosac extract), Water (purified).

Questions or comments? Merx Pharmaceutical Corp., Barrington, IL 1-800-224-4034, M-F 9-4 CST www.releev.com

Shing-RELEEV™
 For the relief of symptoms caused by Shingles and for the treatment of skin infections, minor wounds and other skin irritations.

Even better when used with RELEEV™ INTERNAL SUPPORT

COUPON PRINTED INSIDE BOX

Shing-RELEEV™

Topical Antiseptic Pain Relief & Skin Protectant Spray

- Relief of symptoms associated with Shingles
- Minor wounds
- Other skin irritations

2 fl. oz. (60ml)

We proudly guarantee this product as we do all our fine products:

- RELEEV™ 1-Day Cold Sore Symptom Treatment - 6 ml
- DEEP RELEEV™ Topical Analgesic™
- RELEEV™ CHK-POX Antiseptic Spray™
- RELEEV™ C&F Caps™
- RELEEV™ Internal Support Caps™

COUPON PRINTED INSIDE BOX

Manufactured for:
 Merx Pharmaceutical™ Corp. • Barrington, IL 60010



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U.S. Patent #'s 6,348,503, 6,355,684, 6,946,480
 other patents pending
 NDC# 63287-419-02

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63287-420
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 mg in 1 mL
Allantoin (UNII: 344S277G0Z) (Allantoin - UNII:344S277G0Z)	Allantoin	5 mg in 1 mL
Benzyl Alcohol (UNII: LKG8494WBH) (Benzyl Alcohol - UNII:LKG8494WBH)	Benzyl Alcohol	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0K00R)	
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:63287-420-02	1 in 1 BOX		
1	NDC:63287-420-01	60 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	07/14/2010	

Labeler - Merix Pharmaceutical Corp. (158385687)

Registrant - Topical Pharmaceutiocal Inc. (831530683)

Establishment

Name	Address	ID/FEI	Business Operations
Topical Pharmaceutiocal Inc.		831530683	manufacture(63287-420)