CHIGG AWAY- benzocaine lotion Humco Holding Group, Inc.

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.

Private Label Chigg Away

CVS Itch Relief & Repellent

Drug Facts

Active Ingredient

Benzocaine 5%

Purpose

Anesthetic

Use

Relieves itching and discomfort due to nonpoisonous insect bites such as chiggers (redbugs), mosquitoes, ticks, fleas, no-see-ums, biting flies, fire ants, bees and wasps; summer's itch.

Warnings

For external use only. Keep away from eyes or other mucous membranes. Not for prolonged use. For use on intact skin only. Do not use on children younger than 2 yrs.

When using this product

discontinue use if the condition persists or if a rash or irritation develops and consult a doctor. As with all pesticides/drugs,

keep out of the reach of children.

In case of accidental ingestion, contact physician or poison control center at once.

Directions: As an anti-itch.

- Adults and children 2 yrs. of age and over. Apply topically and rub well as needed.
- Children under 2 yrs. of age: Do not use

As an insect (chigger) repellent: Apply around feet, ankles, waist and to skin under all areas of tight clothing and around all openings in outer clothing. Reapply after heavy perspiration.

Other Information

• **Disposal**: Do not reuse bottle. Rinse thoroughly before discarding.

Inactive Ingredient

Cetyl alcohol. Glycerin, Glyceryl stearate, Hydroxyethyl cellulose, Isopropyl alcohol, Methylparaben,

Petrolatum, Propylparaben, Purified water, Sodium lauryl sulfate, Stearic acid, Sulfur, and Triothanolamine..

CVS



Warning: It is a violation of Federal law to use this product in a manner inconsistent with its labeling. Drug Facts Active ingredient Purpose Benzocaine 5%. Anesthetic Uses relieves itching and discomfort due to nonpoisonous insect bites such as chiggers (redbugs), mosquitoes, ticks, fleas, no-see-ums, biting flies, fire ants, bees and wasps; swimmer's itch. Warnings ■ For external use only. ■ Keep away from eyes or other mucous membranes. ■ Not for prolonged use. ■ For use on intact skin only. Do not use on children younger than 2 yrs. When using this product discontinue use if the condition persists or if rash or irritation develops and consult a doctor. As with all pesticides/drugs, keep out of the reach of children. In case of accidental ingestion, contact a physician or poison control center at once. Directions: As an anti-itch: Adults and children 2 yrs. of age and over: apply topically and rub well as needed. Children under 2 yrs. of age: do not use. ∞ As an insect (chigger) repellent: Apply around feet, ankles, waist and to skin under all areas of tight clothing and around all openings in outer clothing. Reapply after heavy perspiration. ∞ 2 Other information Disposal: Do not reuse bottle Rinse thoroughly before discarding. Inactive ingredients
Cetyl Alcohol, Glycerin, Glyceryl S Stearate, Hydroxyethyl Cellulose, Isopropyl Alcohol, Methylparaben, Petrolatum, Propylparaben, Purified Water, Sodium Lauryl Sulfate, Stearic Acid, Sulfur, Triethanolamine.



Warning: It is a violation of Federal law 6840-01-137-8456 to use this product in a manner inconsistent with its labeling.

Drug Facts

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Directions

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CHIGG AWAY

benzocaine lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0395-9105

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)
BENZOCAINE
50 mg in 1 mL

Inactive Ingredients Ingredient Name Strength CETYL ALCOHOL (UNII: 936JST6JCN) GLYCERIN (UNII: PDC6A3C0OX) GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4) HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: L605B5892V) ISOPROPYL ALCOHOL (UNII: ND2M416302) METHYLPARABEN (UNII: A218C7H19T) PETROLATUM (UNII: 4T6H12BN9U) PROPYLPARABEN (UNII: Z81X2SC1OH) WATER (UNII: 059QF0KOOR)

SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
SULFUR (UNII: 70 FD1KFU70)		
TROLAMINE (UNII: 903K93S3TK)		

# Item Code	Package Description	Marketing Start Date	Marketing End Date
	3 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination oduct	0 1/0 1/20 0 8	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part348	0 1/0 1/20 0 8				

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment			
Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	analysis(0395-9105), manufacture(0395-9105), pack(0395-9105), label(0395-9105)

Revised: 6/2020 Humco Holding Group, Inc.