CHIGG AWAY- benzocaine lotion
Pharma Nobis, LLC

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

CHIGG AWAY

benzocaine lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82645-918
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**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: A2I8C7HI9T)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SULFUR (UNII: 70FD1KFU70)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82645-918-94	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2008	

Labeler - Pharma Nobis, LLC (118564114)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(82645-918) , manufacture(82645-918) , pack(82645-918) , label(82645-918)

Revised: 12/2023 Pharma Nobis, LLC