CHIGG AWAY- benzocaine lotion
Westlake HB Pharma LLC

Chigg Stop

Chigg Away

Drug Facts

Active Ingredient

Benzocaine 5%

Purpose

Anesthetic

Uses

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

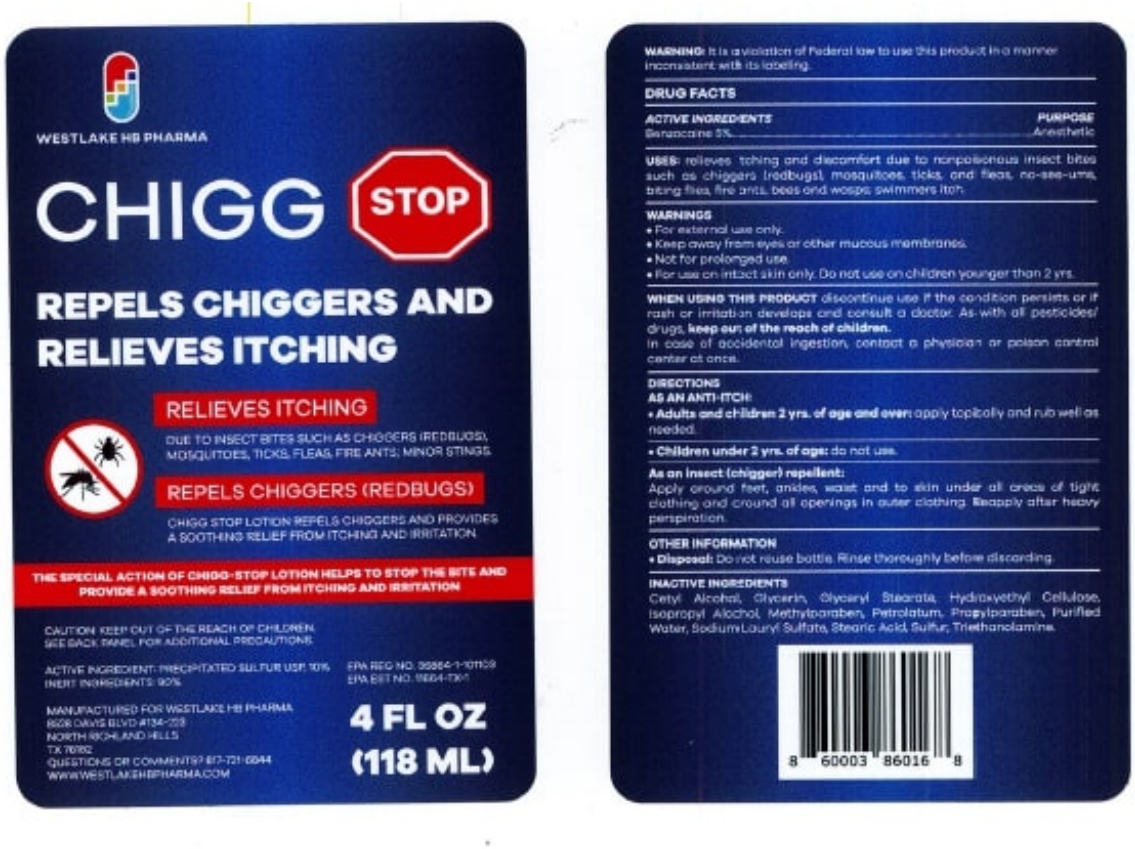
Other Information

- 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, Triethanolamine..

PRINCIPAL DISPLAY PANEL



NDC 86000-857-05

CHIGG

STOP

RELIEVES ITCHING

REPELS CHIGGERS

4 FL OZ

(118 ML)

CHIGG AWAY

benzocaine lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82895-857
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)	
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)	
HYDROXYETHYL CELLULOSE (1500 MPA.S AT 1%) (UNII: L605B5892V)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:82895-857-05	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/23/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/23/2022	

Labeler - Westlake HB Pharma LLC (081176855)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(82895-857) , pack(82895-857) , manufacture(82895-857) , label(82895-857)

Revised: 12/2023

Westlake HB Pharma LLC