ZEP E-2 FOAMING SANITIZING HAND CLEANER- chloroxylenol liquid
Zep 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.

66949-994 E-2 Foaming

Active ingredient

Chloroxylenol 3.0%

Purpose

Antiseptic Hand Wash

Uses

Hand sanitizing to decrease bacteria on the skin.

Warnings

For external use only. Do not usein or around the eyes; if in eyes, rinse promptly and thoroughly with water.

When using this product

- Avoid eye contact.
- If in the eyes, rinse promptly and throoughly with water. Do not swallow.
- If swallowed, do not induce vomiting and if individual is conscious, give large quantities of water to dring and consult a physician immediately.
- Allergy Alert: May cause skin reactions. Symptoms may include: skin reddening, blisters and rash. If a skin reaction occurs, stop use and seek medical help right away.

Stop use and ask a doctor if skin irritation or redness persists for more than 72 hours.

Keep out of reach of children and pets. Children must be supervised in use of this product.

If swallowed, get medical help or contact a Poison Control center immediately.

Directions

- We hands with water.
- Apply liquid Soap.
- Massage soap into hands and wrists, emphasizing back of hands, knuckles and cuticles.
- Rinse hands thoroughly and dry.

Other information

- Store at 20 to 25°C (68 to 77°F).
- Dispose in accordance with all applicable federal, local and state regulations.

Inactive ingredients

Water, Sodium C14-16 Olefin Sulfonate, Triethylene Glycol, Cocamidopropyl Betain, Sodium Xylenesulfonate, PEG-75 Lanolin, Citric Acid

Questions or comments?

Call 1-800-I-BUY-ZEP (1-800-428-9937)

ZEP E-2 FOAMING SANITIZING HAND CLEANER

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-994
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-994-16	2500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/19/2000	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/19/2000	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-994)

Revised: 12/2019 Zep Inc.