AFCO 5508 SANIFECT FOAM-E II- chloroxylenol liquid
Zep Inc.

66949-141 / AF5508 AFCO Sanifect Foam-E II

Active ingredient

Chloroxylenol 1.0% w/w

Purpose

Antibacterial

Uses

- For handwashing to decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

FOR EXTERNAL USE ONLY

When using this product

When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor

Stop use and ask a doctor if if irritation or redness develops and persists.

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact poison control center immediately.

Directions

- Wet hands with water, apply enough product to cover all hand surfaces, rub hands together for at least 30 seconds to work into lather.
- Rinse thoroughly and dry hands completely.

Other information

For Food Processing, Food Service and Professional Use Only.

Inactive ingredients

Water, Ammonium Lauryl Sulfate, Ethanol, Sodium Lauroyl Sarcosinate, Triethylene Glycol, Propylene Glycol, Glycerin, Disodium EDTA, Cocomidopropyl Hydroxysultaine, Citric Acid, Cocamide MIPA.

Questions or comments?

For product or technical information, contact ZEP, INC. Monday to Friday 8 AM to 4 PM EST at 1-877-428-9937 or visit our website at www.zep.com.

AFCO 5508 SANIFECT FOAM-E II

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-141
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	2081980 mg in 208198 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALCOHOL (UNII: 3K9958V90M)	
COCAMIDE MEA (UNII: C80684146D)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
CITRIC ACID (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-141-01	208198 mL in 1 DRUM; Type 0: Not a Combination Product	10/23/2025	
2	NDC:66949-141-02	6000 mL in 1 CASE; Type 0: Not a Combination Product	10/23/2025	
3	NDC:66949-141-03	3785 mL in 1 CASE; Type 0: Not a Combination Product	10/23/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/23/2025	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-141)

Revised: 12/2025 Zep Inc.