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

66949-139 / AF5508 AFCO Sanifect Foam-E II

Active ingredient

Chloroxylenol 1.0% w/w

Purpose

Antiseptic

Uses

- Hand wash to help reduce bacteria on skin that potentially can cause diseases.
- Helps prevent cross contamination by hand contact.
- Recommended for repeat use.

Warnings

For external use only – Harmful if swallowed.

When using this product

When using this product do not use in or near eyes. If eye contact occurs, rinse eyes thoroughly with water. Discontinue use if irritation or redness develops.

Stop use and ask a doctor

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

Keep out of reach of children

Keep out of reach of children. In case of accidental ingestion, seek medical attention or contact poison control center immediately.

Directions

- For one step hand washing and sanitizing, wet hands with water.
- Apply a generous amount of product in palm of hand and work into later.
- Rinse thoroughly with potable water.
- Dry hands completely.

Other information

For Food Processing, Food Service and Professional Use Only.

Inactive ingredients

Water, Ammonium Lauryl Sulfate, Ammonium Laureth Sulfate, Ethyl Alcohol, Propylene Glycol, Cocamide MEA, Glycerin, Disodium EDTA.

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

ethanol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-139
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
AMMONIUM LAURETH-2 SULFATE (UNII: 698O4Z48G6)	
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALCOHOL (UNII: 3K9958V90M)	
COCAMIDE MEA (UNII: C80684146D)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-139-01	208198 mL in 1 DRUM; Type 0: Not a Combination Product	05/05/2025	12/31/2026
2	NDC:66949-139-02	6000 mL in 1 CASE; Type 0: Not a Combination Product	05/05/2025	12/31/2026
3	NDC:66949-139-03	3785 mL in 1 CASE; Type 0: Not a Combination Product	05/05/2025	12/31/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/05/2025	12/31/2026

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-139)

Revised: 12/2025 Zep Inc.