

PREMIUM FOAM ANTIBACTERIAL- chloroxylenol liquid
GOJO Industries, Inc.

GOJO Premium Foam Antibacterial Handwash

Active ingredient

Chloroxylenol 0.3%

Purpose

Antimicrobial

Uses

- Handwash to help decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

Inactive ingredients

Water (Aqua), Alcohol, Ammonium Laureth Sulfate, Ammonium Lauryl Sulfate, Propylene Glycol, Ammonium Xylenesulfonate, Cocamide MEA, Glycerin, Isopropyl Alcohol, Lactic Acid, Retinyl Palmitate, Simmondsia Chinensis (Jojoba) Seed Oil, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Ammonium Sulfate, Fragrance (Parfum), Methylchlorisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)



Drug Facts	
Active Ingredient Chloroxylenol 0.3%.....	Purpose Antimicrobial
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Drug Facts (continued)	
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5362-644-ES-B	

PREMIUM FOAM ANTIBACTERIAL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-085
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.003 mg in 1 mL	
Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			
AMMONIUM LAURETH-2 SULFATE (UNII: 698O4Z48G6)			
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
AMMONIUM XYLENESULFONATE (UNII: 4FZY6L6XCM)			
COCO MONOETHANOLAMIDE (UNII: C80684146D)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL ALCOHOL (UNII: ND2M416302)			
LACTIC ACID (UNII: 33X04XA5AT)			
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)			
JOJOBA OIL (UNII: 724GKU717M)			
EDETATE SODIUM (UNII: MP1J8420LU)			

.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CORN OIL (UNII: 8470G57WFM)	
AMMONIUM SULFATE (UNII: SU46BAM238)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
WATER (UNII: 059QF0K00R)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-085-23	222 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	
2	NDC:21749-085-80	800 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	05/31/2021
3	NDC:21749-085-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	
4	NDC:21749-085-46	1360 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	05/31/2019

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/11/2012	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-085)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-085) , pack(21749-085)

Revised: 7/2025

GOJO Industries, Inc.