

**PROVON FOAMING ANTIMICROBIAL HANDWASH WITH MOISTURIZERS-
chloroxylenol liquid
GOJO Industries, Inc.**

PROVON Foaming Antimicrobial Handwash with Moisturizers

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), Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)

5286

NDC 21749-090-67
Brought to you by GOJO
Llega a usted gracias a GOJO

PROVON[®]
BRAND

Foaming Antimicrobial Handwash
with Moisturizers
Jabón en Espuma Antibacterial
con Humectantes

gojo[®]

Distributed by, Distribuido por:
GOJO Industries, Inc.
Akron, OH 44309
800-321-8647 • 330-255-6000
www.GOJO.com
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Made in U.S.A., Hecho en los E.U.U.

HAND WASH
JABÓN PARA MANOS

5286-645-ES

2 L (67 US/ÉU FL OZ)

Drug Facts		Drug Facts (continued)	
Active Ingredient	Purpose	Directions	
Chloroxylenol 0.3%	Antimicrobial	<ul style="list-style-type: none"> Wet hands Apply product and thoroughly cover hands with lather Rinse well and dry hands completely 	
Uses		Inactive Ingredients	
<ul style="list-style-type: none"> Handwash to help decrease bacteria on the skin Recommended for repeated use 		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), Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)	
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.			

PROVON FOAMING ANTIMICROBIAL HANDWASH WITH MOISTURIZERS

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-090
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
WATER (UNII: 059QF0KO0R)	

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)
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-090-22	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/13/2003	11/26/2024
2	NDC:21749-090-42	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/13/2003	
3	NDC:21749-090-67	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	07/13/2003	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/13/2003	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-090)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-090) , pack(21749-090)

