

GOJO ANTIMICROBIAL FOAM HANDWASH WITH PCMX- chloroxylenol liquid
GOJO Industries, Inc.

GOJO Antimicrobial Foam Handwash with PCMX

Active ingredient

Chloroxylenol 0.5%

Purpose

Antimicrobial

Use

- 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 product and thoroughly cover hands with lather.
- Rinse well and dry hands completely.

Inactive ingredients

Water (Aqua), Alcohol, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Sodium Sulfite, Tetrasodium EDTA, Sodium Sulfate, Fragrance (Parfum), Methylparaben, Propylparaben, Green 3 (CI 42053), Red 33 (CI 17200)

1340

**ANTIMICROBIAL FOAM
HANDWASH WITH PCMX**
**JABÓN DE MANOS
ANTIBACTERIAL EN ESPUMA
CON CLOROXILENOL**

Distributed by, Distribuido por:
GOJO Industries, Inc., Akron, OH 44309
Tel: 800-321-0647 • 330-255-6000
www.GOJO.com
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Made in U.S.A., Hecho en los E.E.U.U.

700 mL (23.6 US/ÉU FL OZ)

1340-640-ES

Drug Facts

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Chloroxylenol 0.5%.....	Antimicrobial

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Drug Facts (continued)

Directions

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GOJO ANTIMICROBIAL FOAM HANDWASH WITH PCMX

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-521
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.005 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
LAURIC ACID (UNII: 1160N9NU9U)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
LACTIC ACID (UNII: 33X04XA5AT)	
POLOXAMER 124 (UNII: 1S66E28KXA)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-521-53	535 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2014	12/02/2024
2	NDC:21749-521-97	700 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2014	07/31/2021
3	NDC:21749-521-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2014	06/30/2023
4	NDC:21749-521-90	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/30/2014	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-521)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-521) , pack(21749-521)

Revised: 12/2024

GOJO Industries, Inc.