MICRELL ANTIBACTERIAL FOAM HANDWASH - chloroxylenol liquid
GOJO Industries, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

MICRELL Antibacterial Foam Handwash

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)

Package/Label Principal Display Panel

MICRELL Antibacterial Foam Handwash

Brought to you by GOJO
Distributed by

GOJO Industries, Inc. Akron, OH 44309

Questions? 800-321-9647 • 330-255-6000 www.GOJO.com

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MICRELL ANTIBACTERIAL FOAM HANDWASH

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-095
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:21749-095-22	200 mL in 1 BOTTLE		
2	NDC:21749-095-53	535 mL in 1 BOTTLE		
3	NDC:21749-095-97	700 mL in 1 BOTTLE		
4	NDC:21749-095-89	1200 mL in 1 BOTTLE		
5	NDC:21749-095-90	1250 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/23/2006	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-095)

Revised: 6/2006 GOJO Industries, Inc.