MICRELL ANTIBACTERIAL LTN SP- chloroxylenol liquid
GOJO INDUSTRIES, INC.

MICRELL Antibacterial Lotion Soap

Active ingredient

Chloroxylenol 0.3%

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

Inactive ingredients

Water (Aqua), Coconut Acid, Oleic Acid, Sodium Sulfate, Ethanolamine, Cocamide MEA, Coco-Betaine, Propylene Glycol, Retinyl Palmitate, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Hydroxypropyl Methylcellulose, Fragrance (Parfum)

Drug Facts

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Questions or comments?

Call **1-800-321-9647** Monday through Friday 8:00 AM to 5:00 PM

MICRELL ANTIBACTERIAL LTN SP

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-975
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)	
COCONUT ACID (UNII: 40U37V505D)	
OLEIC ACID (UNII: 2UMI9U37CP)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
COCO-BETAINE (UNII: 03DH2IZ3FY)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN OIL (UNII: 8470G57WFM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-975-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/1998	12/09/2024
2	NDC:21749-975-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/1998	
3	NDC:21749-975-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/1998	
4	NDC:21749-975-50	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/1998	12/09/2024
5	NDC:21749-975-80	800 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/1998	03/31/2023
6	NDC:21749-975-10	1000 mL in 1 BAG; Type 0: Not a Combination Product	07/30/1998	
7	NDC:21749-975-33	1000 mL in 1 PACKAGE; Type 0: Not a Combination Product	07/30/1998	12/09/2024
8	NDC:21749-975-20	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/1998	12/31/2026
9	NDC:21749-	3784 mL in 1 BOTTLE; Type 0: Not a Combination	07/30/1998	

9	975-37	Product	07/30/1998	
10	NDC:21749-975-52	196841 mL in 1 DRUM; Type 0: Not a Combination Product	07/30/1998	12/09/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/30/1998	

Labeler - GOJO INDUSTRIES, INC. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-975)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-975) , pack(21749-975)

Establishment

Name	Address	ID/FEI	Business Operations
Travis Association for the Blind		026032268	label(21749-975) , pack(21749-975)

Revised: 12/2024 GOJO INDUSTRIES, INC.