

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)

Brought to you by GOJO
Llega a usted gracias a GOJO

NDC 21749-095-97

gojo 1310-03



ANTIBACTERIAL
FOAM HANDWASH
LAVADO DE MANOS
ANTIBACTERIANO
EN ESPUMA

Distributed by; Distribuido por:
GOJO Industries, Inc.
Akron, OH 44309
Questions? ¿Preguntas?
800-321-9647 • 330-255-6100
www.GOJO.com
©2011, GOJO Industries, Inc.
All rights reserved.
Todos los derechos reservados.
Made in U.S.A.
Hecho en los E.E.U.U.

1310-640-ES



700 mL (23.6 US/ÉU FL OZ)

Drug Facts

| | |
|--|---------------------------------|
| Active ingredient Chloroxylenol 0.5% | 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 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)

U.S. Pat 7,851,424
SDA-36-1301

H
I
N
G
E

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; Type 0: Not a Combination Product | 06/23/2006 | |
| 2 | NDC:21749-095-53 | 535 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package | 06/23/2006 | |
| 3 | NDC:21749-095-97 | 700 mL in 1 BOTTLE; Type 0: Not a Combination Product | 06/23/2006 | |
| 4 | NDC:21749-095-89 | 1200 mL in 1 BOTTLE; Type 0: Not a Combination Product | 06/23/2006 | |
| 5 | NDC:21749-095-90 | 1250 mL in 1 BOTTLE; Type 0: Not a Combination Product | 06/23/2006 | |

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) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-----------------------|---------|-----------|------------------------------------|
| GOJO Industries, Inc. | | 088312414 | label(21749-095) , pack(21749-095) |