

PRIMORY ANTIMICROBIAL FOAM- chloroxylenol liquid
GOJO Industries, Inc.

PRIMORY Antimicrobial Foam Soap

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

■ P6162-04



Made in USA for:
 Hecho en los E.E.U.U. por:
Primorance, Inc.
 Akron, OH 44309
 1-800-321-9647
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ANTIMICROBIAL FOAM SOAP
JABÓN ANTIMICROBIANO
EN ESPUMA

1.25 L (42 US FL OZ)

6162-640-PMY-F

Drug Facts

| | |
|--------------------------|----------------|
| Active Ingredient | Purpose |
| Chloroxylenol 0.3%..... | 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

Drug Facts (continued)

Directions

- Wet hands
- Apply product and thoroughly cover hands with 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)

5362-644-ES-E

| | | | |
|--|----------------|---------------------------|-----------------|
| PRIMORY ANTIMICROBIAL FOAM | | | |
| chloroxylenol liquid | | | |
| Product Information | | | |
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:21749-555 |
| 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:21749-555-89 | 1200 mL in 1 PACKAGE; Type 0: Not a Combination Product | 09/09/2004 | 11/26/2024 |
| 2 | NDC:21749-555-90 | 1250 mL in 1 PACKAGE; Type 0: Not a Combination Product | 09/09/2004 | |
| 3 | NDC:21749-555-43 | 1500 mL in 1 PACKAGE; Type 0: Not a Combination Product | 09/09/2004 | 11/26/2024 |
| 4 | NDC:21749-555-67 | 2000 mL in 1 PACKAGE; Type 0: Not a Combination Product | 09/09/2004 | 11/26/2024 |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | 505G(a)(3) | 09/09/2004 | |

Labeler - GOJO Industries, Inc. (004162038)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-----------------------|---------|-----------|------------------------|
| GOJO Industries, Inc. | | 036424534 | manufacture(21749-555) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-----------------------|---------|-----------|------------------------------------|
| GOJO Industries, Inc. | | 088312414 | label(21749-555) , pack(21749-555) |

Revised: 11/2024

GOJO Industries, Inc.