

PRO-LINK PROFORMANCE FOAMING ANTIMICROBIAL SP- chloroxylenol liquid
Pro-Link, Inc.

PRO•LINK® ProFormance™ Foaming Antimicrobial 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 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), Methylchlorisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)



Foaming Antimicrobial Soap

Jabón Antimicrobiano Espumante

Distributed by/Distribuido por Pro-Link, Inc. 1-800-745-4657
 Canton, MA 02021 USA www.prolinkhq.com
 Made in U.S.A./Hecho en los E.E.U.U.



Drug Facts

Active Ingredient **Purpose**
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Datos Farmacológicos

Ingredientes activo **Propósito**
 Cloroxileno 0,3%.....Antimicrobiano

Usos • Lavado de manos empleado para disminuir la cantidad de bacterias en la piel • Recomendado para uso reiterado

Advertencias Sólo para uso externo

Al utilizar este producto, evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamente los ojos con agua.

Dejar de usar el producto y consultar a un médico si aparece y persiste una irritación o erupción cutánea

Mantener fuera del alcance de los niños. En caso de ingestión, de inmediato acudir a un médico o ponerse en contacto con un centro para el control de tóxicos.

Modo De Uso • Moje las manos
 • Aplique el producto hasta cubrir completamente sus manos con la espuma • Enjuague bien y seque las manos completamente

Ingredientes Inactivos Agua, Alcohol etílico, Lauret sulfato de amonio, Laurilsulfato de amonio, Propilenglicol, Xilenosulfonato de amonio, MEA cocamida, Glicerina, Alcohol isopropílico, Acido láctico, Palmitato de retinilo, Aceite de Simmondsia Chinensis, EDTA tetrasódico, Acetato de tocoferilo, Aceite de maíz (Zea Mays), Sulfato de amonio, Fragancia, Metilcloroisotiazolinona, Metilisotiazolinona, Rojo 4, Amarillo 6

5162-640-PD3-B

RBL4

MMH250

1250 mL (42 US/ÉU FL OZ)

PRO-LINK PROFORMANCE FOAMING ANTIMICROBIAL SP

chloroxyleneol liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:66908-083 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|----------|
| CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - | CHLOROXYLENOL | 0.3 mg |

UNII:0F32U78V2Q)

CHLOROXALLENOL

in 100 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:66908-083-42 | 1250 mL in 1 PACKAGE; Type 0: Not a Combination Product | 08/28/2023 | |
| 2 | NDC:66908-083-67 | 2000 mL in 1 PACKAGE; Type 0: Not a Combination Product | 08/28/2023 | 12/31/2025 |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | 505G(a)(3) | 08/28/2023 | |

Labeler - Pro-Link, Inc. (144650637)

Revised: 1/2026

Pro-Link, Inc.