PRIMORY ANTIMICROBIAL FOAM- 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.

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

P6162-04

Primory®

Made in USA for:
Hecho en los E.E.U.U. por:
Primorance, Inc.
Akron, OH 44309
1-800-321-9647

0 73852 02679 5

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

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Stop use and ask a doctor if irritation or rash appears and lasts

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Directions

- Wet hands
- Apply product and thoroughly cover hands with lather
- Rinse well and dry hands completely

Drug Facts (continued)

Directions

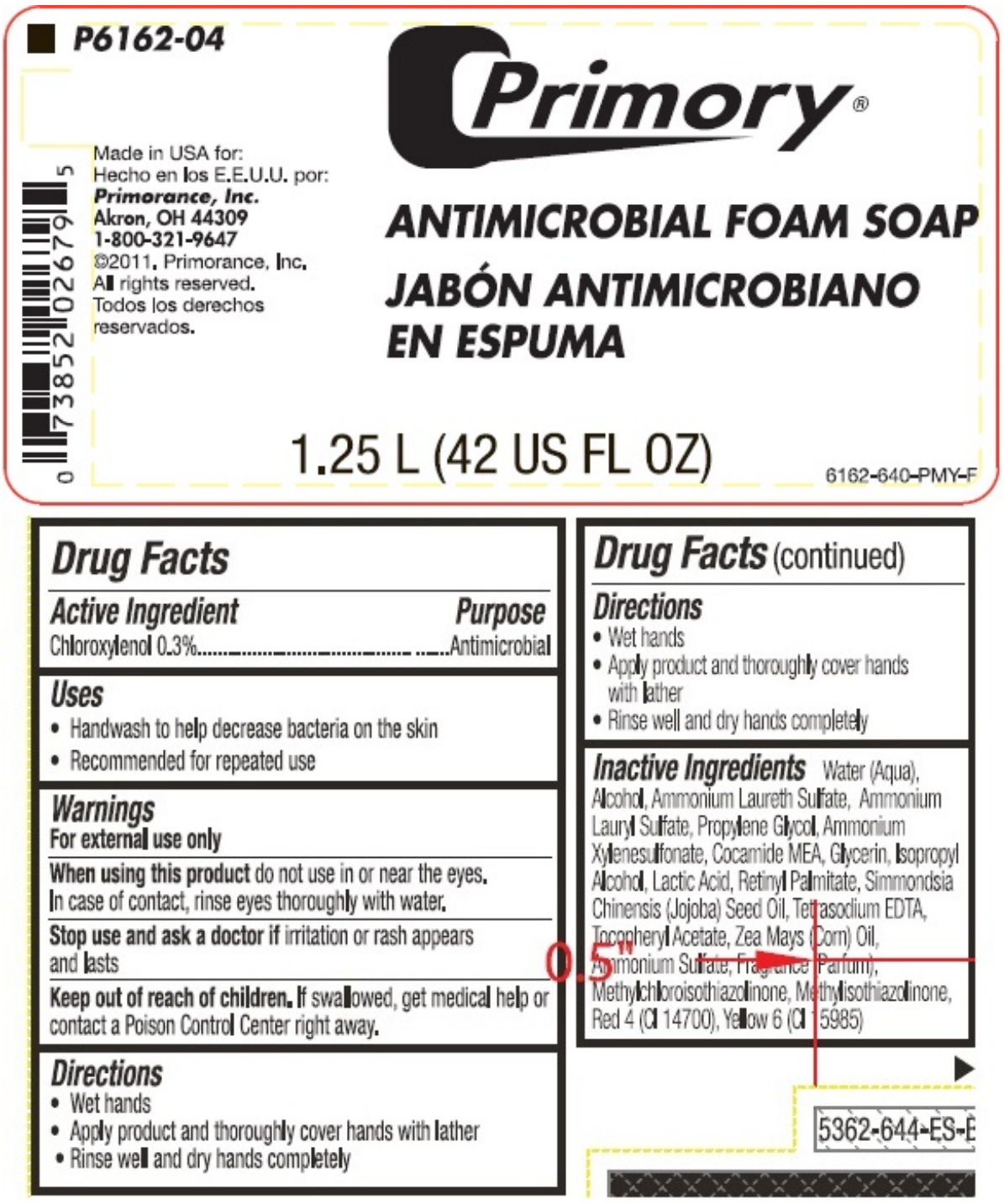
- Wet hands
- Apply product and thoroughly cover hands with lather
- Rinse well and dry hands completely

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)

0.5"

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	
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	
4	NDC:21749-555-67	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2004	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	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: 6/2023

GOJO Industries, Inc.