

LUXURY FOAM ANTIBACTERIAL- 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.

GOJO Luxury Foam Antibacterial Handwash

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)

5562



Distributed by/Distribuido por:
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 Akron, OH 44309
 Questions? ¿Preguntas?
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**LUXURY FOAM
 ANTIBACTERIAL HANDWASH**

**ESPUMA DE LUJO
 ANTIBACTERIANA
 PARA LAVADO DE MANOS**

**For general handwashing and germ killing
 Para el lavado de las manos en general y para
 matar gérmenes**



5562-641-ES-F

800 mL 27 US/ÉU FL OZ NET CONT./CONT. NET 800 mL

Drug Facts

Active Ingredient	Purpose
Chloroxylenol 0.3%	Antimicrobial

Uses

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Warnings
 For external use only

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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 5985)

5362-644-ES-E

LUXURY FOAM ANTIBACTERIAL

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-080
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:21749-080-53	535 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	
2	NDC:21749-080-80	800 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/11/2012	
3	NDC:21749-080-42	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	
4	NDC:21749-080-43	1500 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	
5	NDC:21749-080-67	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/11/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/11/2012	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	MANUFACTURE(21749-080)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-080) , pack(21749-080)

Revised: 12/2019

GOJO Industries, Inc.