PURELL HEALTHY SP 0.5PCT PCMX ANTIMICROBIAL FOAM-
chloroxylenol liquid
GOJO Industries, Inc.

PURELL HEALTHY SOAP 0.5% PCMX Antimicrobial Foam

Active ingredient

Chloroxylenol 0.5%

Purpose

Antimicrobial

Use

- Handwash to help reduce 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, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Tetrasodium EDTA, Fragrance (Parfum), Methylparaben, Propylparaben

PURELL HEALTHY SP 0.5PCT PCMX ANTIMICROBIAL FOAM

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-622
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.5 mg in 100 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)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-622-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/23/2022	
2	NDC:21749-622-90	1250 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/23/2022	
3	NDC:21749-622-17	515 mL in 1 PACKAGE; Type 0: Not a Combination Product	11/14/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/23/2022	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-622) , pack(21749-622) , label(21749-622)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	pack(21749-622) , label(21749-622)

Revised: 12/2024 GOJO Industries, Inc.