PROVON FOAMING ANTIMICROBIAL HANDWASH WITH PCMX-chloroxylenol liquid
GOJO Industries, Inc.

PROVON Foaming Antimicrobial Handwash with PCMX

Active ingredient

Chloroxylenol 0.5%

Purpose

Antimicrobial

Use

- 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

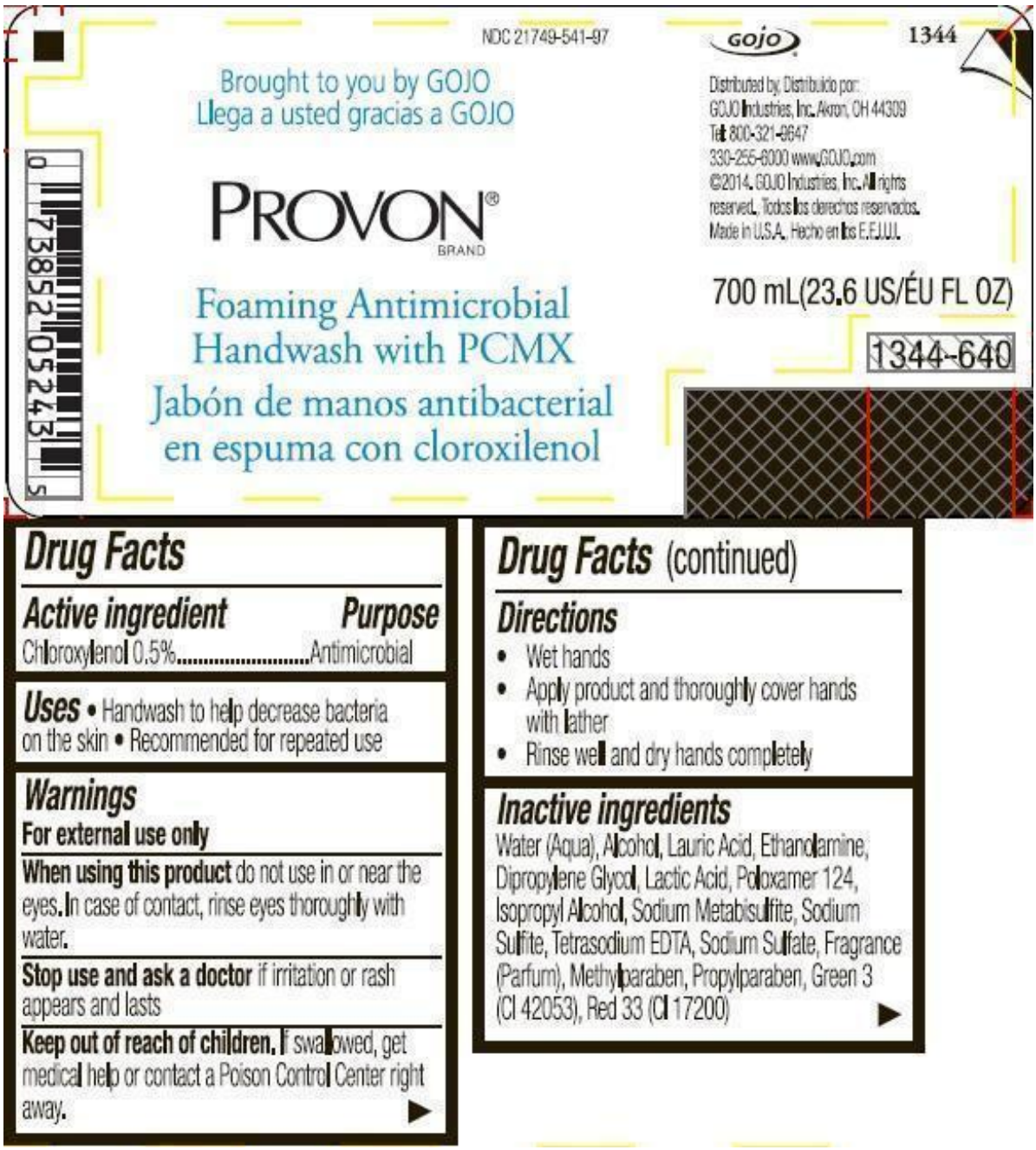
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

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

Inactive ingredients

Water (Aqua), Alcohol, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Sodium Sulfite, Tetrasodium EDTA, Sodium Sulfate, Fragrance (Parfum), Methylparaben, Propylparaben, Green 3 (CI 42053), Red 33 (CI 17200)



PROVON FOAMING ANTIMICROBIAL HANDWASH WITH PCMX

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-541
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.005 mg in 1 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)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-541-53	535 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	04/30/2021
2	NDC:21749-541-97	700 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	
3	NDC:21749-541-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	
4	NDC:21749-541-90	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	
5	NDC:21749-541-20	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	12/02/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/01/2013	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-541)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-541) , pack(21749-541)

Revised: 12/2024

GOJO Industries, Inc.