PROVON MEDICATED FOAM HANDWASH WITH ADVANCED MOISTURIZERS AND TRICLOSAN - triclosan 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.

PROVON Medicated Foam Handwash with Advanced Moisturizers and Triclosan

Active ingredient

Triclosan 0.3%

Purpose

Antimicrobial

Use

- Handwash to help decrease bacteria on the skin before and after contact with a person under medical care or treatment
- 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

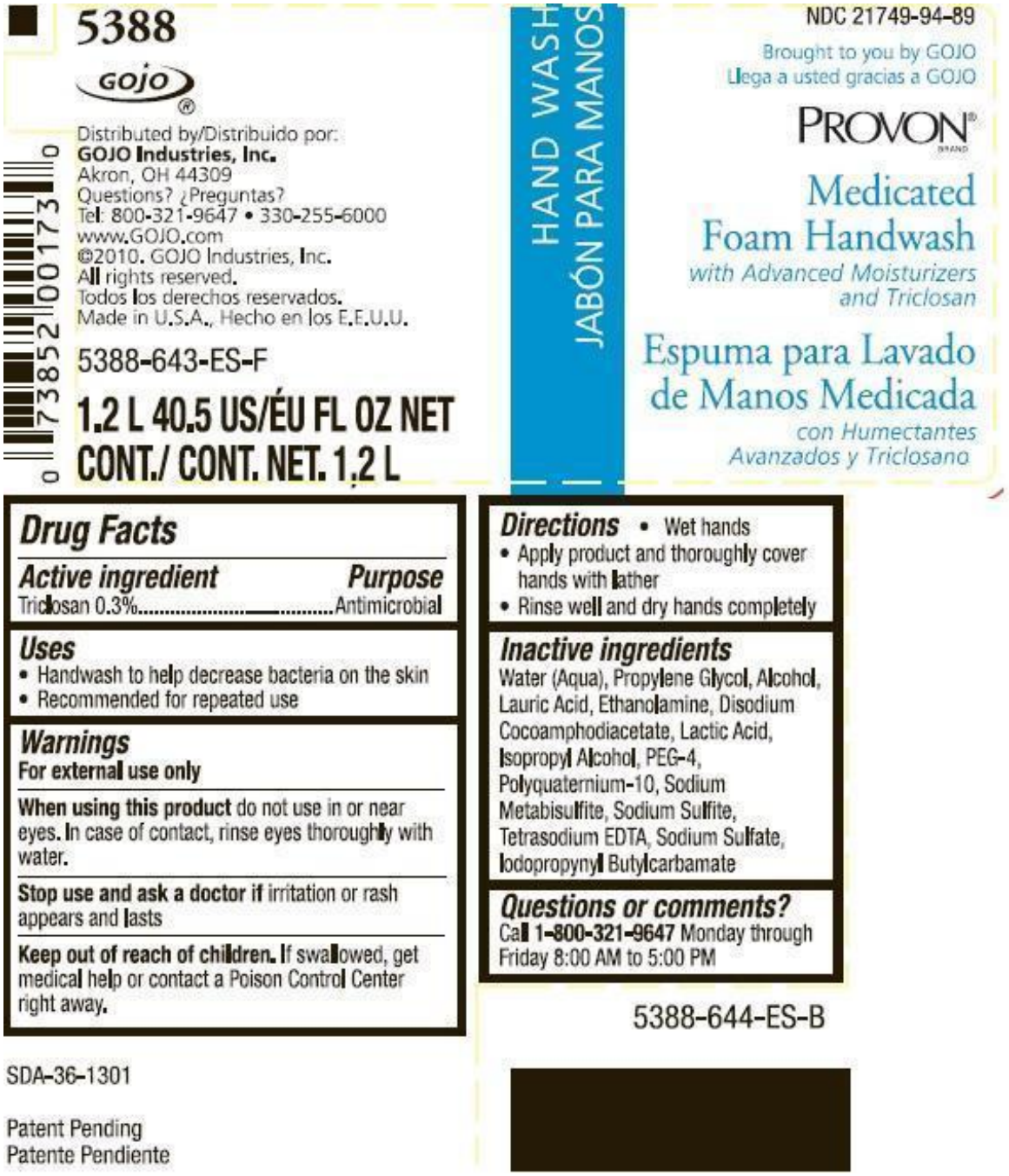
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

Inactive ingredients

Water (Aqua), Propylene Glycol, Alcohol, Lauric Acid, Ethanolamine, Disodium Cocoamphodiacetate, Lactic Acid, Isopropyl Alcohol, PEG-4, Polyquaternium-10, Sodium Metabisulfite, Sodium Sulfite, Tetrasodium EDTA, Sodium Sulfate, Iodopropynyl Butylcarbamate



PROVON MEDICATED FOAM HANDWASH WITH ADVANCED MOISTURIZERS AND TRICLOSAN

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-094
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.003 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
LAURIC ACID (UNII: 1160N9NU9U)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
LACTIC ACID (UNII: 33X04XA5AT)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)	
POLYQUATERNIUM-10 (30000 MPA.S AT 2%) (UNII: C7RDC8Y4JS)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-094-89	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2013	
2	NDC:21749-094-90	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2013	
3	NDC:21749-094-97	700 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2013	
4	NDC:21749-094-53	535 mL in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/01/2013	12/01/2021

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-094)

Revised: 11/2018 GOJO Industries, Inc.