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.

Active ingredient

Triclosan 0.3%

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

Inactive ingredients

Water (Aqua), Coconut Acid, Sodium Sulfate, Oleic Acid, Ethanolamine, Aloe Barbadensis Leaf Juice, Cocamide MEA, Coco-Betaine, Hydrolyzed Vegetable Protein, Propylene Glycol, Retinyl Palmitate, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Hydroxypropyl Methylcellulose, Methylchloroisothiazolinone, Methylisothiazolinone

TRICLOSAN

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21749-415
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)	
COCONUT ACID (UNII: 40U37V505D)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
OLEIC ACID (UNII: 2UMI9U37CP)	

MONOETHANOLAMINE (UNII: 5KV86114PT)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
COCO-BETAINE (UNII: 03DH2IZ3FY)	
Propylene Glycol (UNII: 6DC9Q167V3)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN OIL (UNII: 8470G57WFM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
Methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
Methylisothiazolinone (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-415-10	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/10/2010	
2	NDC:21749-415-20	2000 mL in 1 BOX; Type 0: Not a Combination Product	03/10/2010	
3	NDC:21749-415-16	473 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/10/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/10/2010	12/01/2021

Labeler - GOJO Industries, Inc. (004162038)

Registrant - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-415)

Revised: 10/2016 GOJO Industries, Inc.