PRO ADVANTAGE ANTIBACTERIAL - triclosan liquid
NDC National Distribution & Contracting, 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.

Pro Advantage Antibacterial Liquid Soap

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Active Ingredient

Triclosan, 0.3%

Purpose

Antiseptic Handwash

Use

Handwash to help reduce bacteria that potentially can cause disease.

Warnings

• For external use only

Ask a doctor before use if you have

• Deep wounds, animals bites, or serious burns.

When using this product

• Avoid contact with eyes. If this occurs, rinse thoroughly with water.

Stop use and ask a doctor if

• Irritation, itching or redness develops. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

• If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Wet hands, apply soap, lather for 30 seconds, and rinse hands thoroughly.

Inactive Ingredients

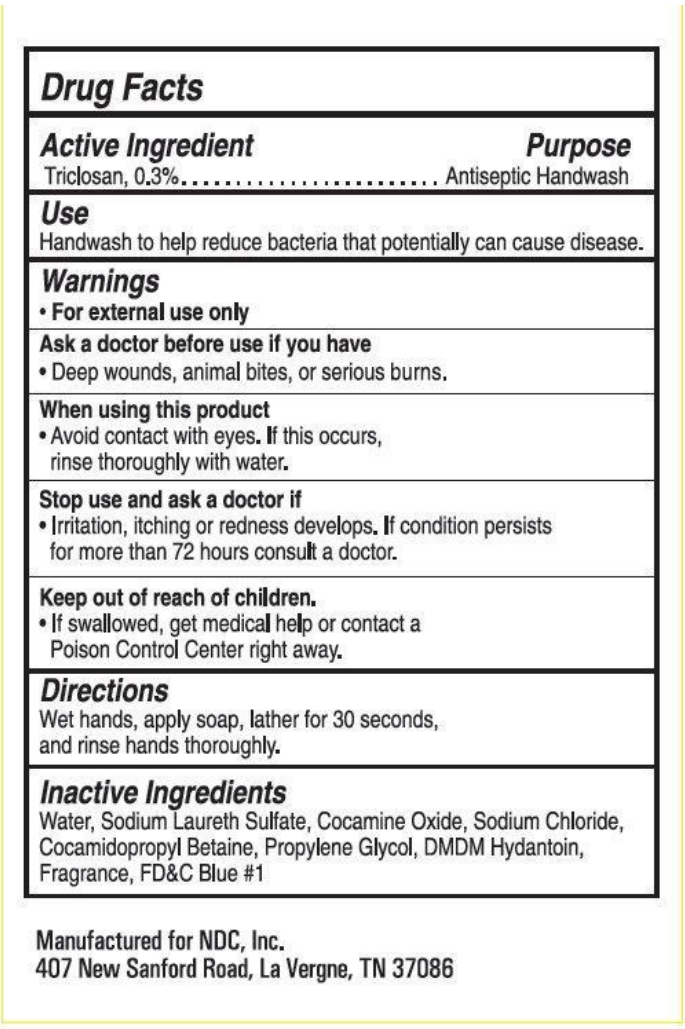
Water, Sodium Laureth Sulfate, Cocamine Oxide, Sodium Chloride, Cocamidopropyl Betaine, Propylene Glycol, DMDM Hydantoin, Fragrance, FDandC Blue #1

REF: P778108 NDC 43128-108-02

Made in China
www.ProAdvantagebyNDC.com

Manufactured for NDC, Inc.
407 New Sanford Road, La Vergne, TN 37086

Product Labels



PRO ADVANTAGE ANTIBACTERIAL

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43128-108
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43128-108-02	1 in 1 PACKAGE		
1		237 mL in 1 BOTTLE, PUMP		
2	NDC:43128-108-03	296 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/08/2013	

Labeler - NDC National Distribution & Contracting, Inc. (009831413)

Revised: 4/2013 NDC National Distribution & Contracting, Inc.