BACTI-FOAM- triclosan solution
Ecolab 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.

Drug Facts

Active Ingredient

Triclosan, 0.3%

Purpose

Healthcare personnel handwash

Uses

- healthcare personnel handwash

Warnings

- **For external use only**

When using this product

- discontinue use if irritation and redness develop

Stop use and ask Doctor if

- skin irritation and redness persist for more than 72 hours

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

Directions

- wet skin and apply a small amount on hands and forearms
- scrub well, rinse thoroughly and dry

Other information

- for emergency medical information in USA and Canada, call 1.800.328.0026
- for emergency medical information worldwide, cal 1.651.222.5352 (in the USA)

Inactive ingredients water, potassium cocoate, SD alcohol 40-B, glycerin, potassium stearate, tetrasodium EDTA, cocamidopropyl PG-dimonium chloride phosphate, cocamine oxide, fragrance, methylparaben, tocopheryl acetate, citric acid, potassium hydroxide, isopropyl alcohol, propylparaben, aloe barbadensis leaf juice, FDC blue 1, FDC yellow 5

Questions? call **1.866.781.8787**

Principal Display Panel and Representative Label

ECOLAB®

NDC 47593-169-56

DIN 02237409

BACTI-FOAM®

Antimicrobial Foam Hand Soap

Healthcare Peronnel Handwash

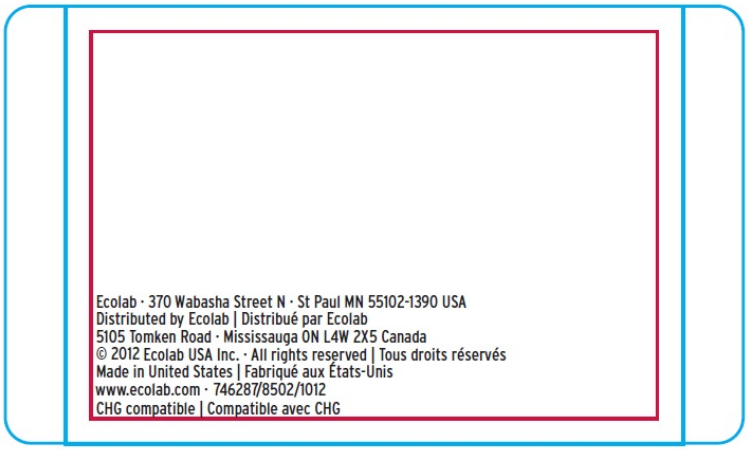
Active Ingredient: 0.3% Triclosan

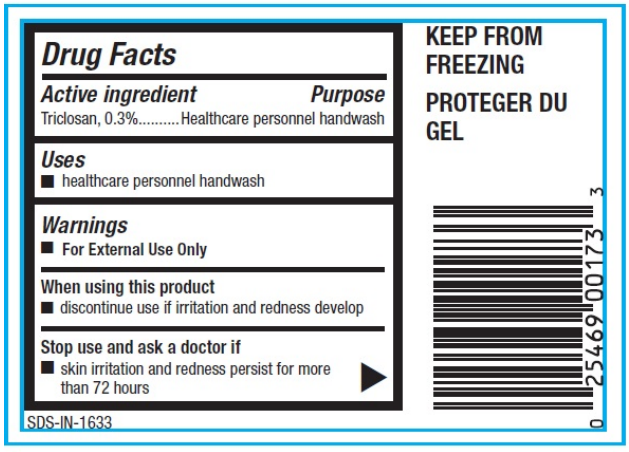
Nets Contents:

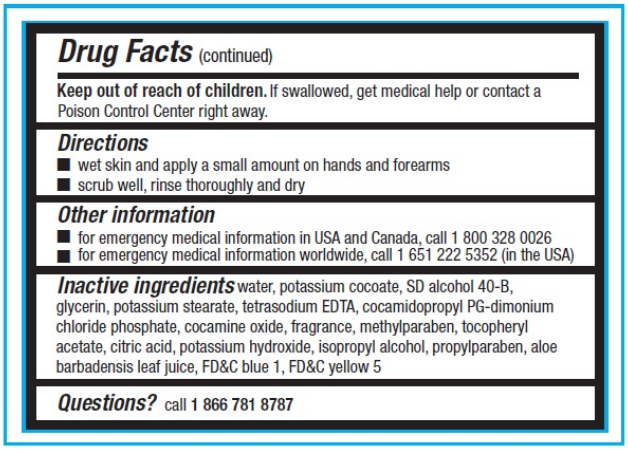
1200 mL (40.6 fl oz)

6082565

747795/8502/1012







BACTI-FOAM

triclosan solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-169
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)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM STEARATE (UNII: 17V812XK50)	
EDETATE SODIUM (UNII: MP1J8420LU)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-169-38	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/08/2012	
2	NDC:47593-169-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/13/2013	
3	NDC:47593-169-56	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/30/2009	
4	NDC:47593-169-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/30/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/11/1997	

Labeler - Ecolab Inc. (006154611)

Revised: 10/2016 Ecolab Inc.