

**ANTIBACTERIAL WITH LIGHT MOISTURIZERS- triclosan soap
BJWC**

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 Box - Back Label

Active Ingredient

Triclosan 0.115%

Purpose

- Antibacterial.

Warnings

For external use only.

When using this product

Avoid contact with eyes. If contact occurs, rinse with water.

Stop using this product and ask doctor if

Irritation or redness develops and lasts.

Keep out of reach of children

In case of accidental ingestions, get medical help or contact a Poison Control Center immediately.

Questions/Comments?

1-800-934-1204

Directions

Apply to wet hands, lather and rinse thoroughly.

Uses

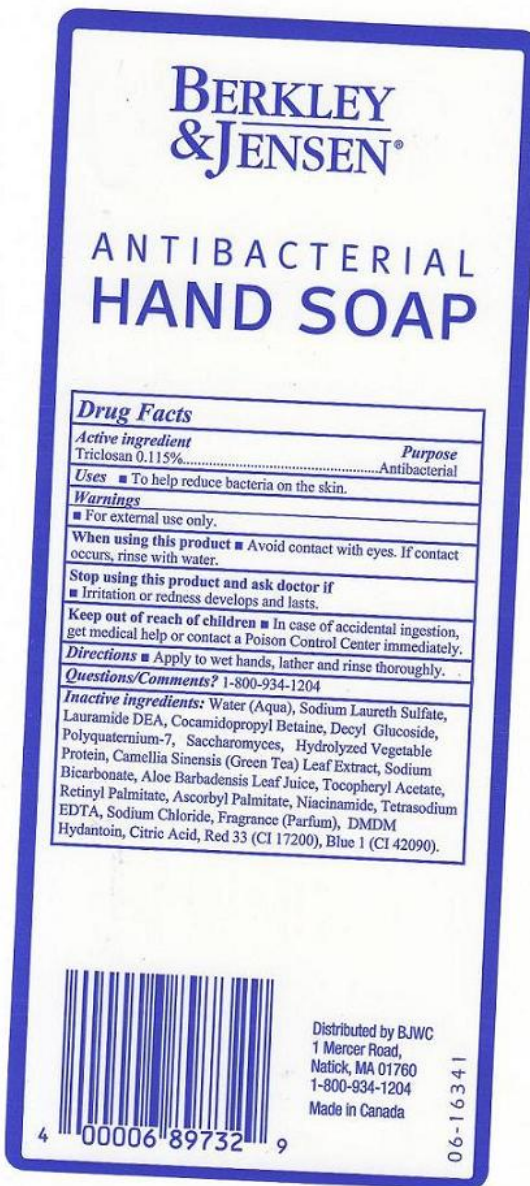
To help reduce bacteria on the skin

Inactive Ingredients

Water (Aqua), Sodium Laureth Sulfate, Lauramide DEA, Cocamidopropyl Betaine, Decyl Glucoside, Polyquaternium-7, Saccharomyces, Hydrolyzed Vegetable Protein, Camellia Sinensis (Green Tea) Leaf Extract, Sodium Bicarbonate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate, Retinyl Palmitate, Ascorbyl Palmitate, Niacinamide, Tetrasodium EDTA, Sodium Chloride, Fragrance (Parfum), DMDM Hydantoin, Citric Acid, Red 33 (CI 17200), Blue 1 (CI 42090).

Package Front and Back Labels

bj1g.jpg



ANTIBACTERIAL WITH LIGHT MOISTURIZERS

triclosan soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68 391-141
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.115 mL in 100 mL
Inactive Ingredients		
Ingredient Name		Strength
WATER (UNII: 059QF0K00R)		
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)		
LAURIC DIETHANOLAMIDE (UNII: I29I2VHG38)		
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)		
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600 KD) (UNII: 0L414VCS5Y)		
SACCHAROMYCES CEREVISIAE (UNII: 978D8U419H)		
GREEN TEA LEAF (UNII: W2ZU1RY8B0)		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)		
ASCORBYL PALMITATE (UNII: QN83US2B0N)		
NIACINAMIDE (UNII: 25X51I8RD4)		
EDETATE SODIUM (UNII: MP1J8420LU)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
DMDM HYDANTOIN (UNII: BYR0546TOW)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
D&C RED NO. 33 (UNII: 9DBA0SBB0L)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68391-141-50	3780 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/23/2010	

Labeler - BJWC (159082692)