SMITH AND JOHNSON ANTIBACTERIAL - triclosan soap Allied International Corp

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.

Smith & Johnson Liquid Soap Antibacterial

Drug Facts

Active Ingredient

Triclosan 0.115%

Purpose

Antiseptic

USES: For hand washing to decrease bacteria on the skin.

WARNING:

For external use only.

When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Keep out of reach of children.

If swallowed, contact a poison control center immediately and consult a physician if necessary.

DIRECTIONS

- Wet hands
- Apply palmful to hands
- Scrub thoroughly
- Rinse

WARNING: AS WITH ANY SOAP, AVOID CONTACT WITH EYES, IN CASE OF CONTACT, FLUSH WITH WATER.

The Smith and Johnson Liquid Soap is formulated to clean dirt and kill germs, an ideal high quality soap for the entire family.

INGREDIENTS: Water, Sodium Polyoxyethylene Fatty Alcohol Sulfate, Coconut Oil Diethanolamide, Citric Acid, Kathon CG, Fragrance, Sodium Chloride, FD and C Yellow No.5, FD and C Red No.40.

ITEM Number: 20063 / 4492

MADE IN CHINA FOR

ALLIED INTERNATIONAL CORP.

Glen Burnie, MD 21060

www.alliedint.com

Smith and Johnson

Liquid Soap

Antibacterial

16.9 FL.OZ (500ml)

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SMITH AND JOHNSON ANTIBACTERIAL

triclosan soap

Product Information

Product Type HUMAN OTC DRUG NDC:52546-110 Item Code (Source)

Route of Administration **TOPICAL**

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X) TRICLOSAN 0.115 mL in 100 mL

Inactive Ingredients Strength **Ingredient Name** WATER (UNII: 059QF0KO0R) SODIUM LAURETH-2 SULFATE (UNII: ZZQ59TY3KG) COCONUT OIL (UNII: Q9L0O73W7L) **DIETHANOLAMINE** (UNII: AZE05TDV2V) CITRIC ACID MONOHYDRATE (UNII: 2968 PHW8 QP) **SODIUM CHLORIDE** (UNII: 451W47IQ8X) FD&C YELLOW NO. 5 (UNII: I753WB2F1M) FD&C RED NO. 40 (UNII: WZB9127XOA)

I	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:52546-110-00	500 mL in 1 BOTTLE					

Marketing Info	rketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333E	05/19/2011				

Labeler - Allied International Corp (004001780)

Revised: 5/2011 Allied International Corp