SMITH AND JOHNSON ANTIBACTERIAL- triclosan liquid
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 and Johnson Antibacterial Liquid Soap

Active ingredient

Triclosan 0.115 percent

Uses

For handwashing to decrease bacteria on the skin.

Warnings

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

Inactive Ingredients

Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Cocamide DEA, Sodium Chloride, Citric Acid, Fragrance, DMDM Hydantoin, Tetrasodium EDTA

Designed in USA

Made in China for

Allied International Corp of VA

PO Box 858 Ashburn, VA 20146 USA

www.alliedint.com

Keep out of reach of children.

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

Purpose:

Antiseptic

Smith and Johnson

New

Liquid Soap

Antibacterial

16 FL OZ (470 ML)

SMITH AND JOHNSON ANTIBACTERIAL

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52546-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	115 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	

DMDM HYDANTOIN (UNII: BYR0546TOW)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52546-100-16	470 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/21/2010	09/28/2016
2	NDC:52546-100-40	1180 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/21/2010	09/28/2016

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/21/2010	

Labeler - Allied International Corp (004001780)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangsu Longliqi Bioscience Co., Ltd.		421547530	manufacture(52546-100)

Revised: 9/2016 Allied International Corp