WHITE LAVENDER AND MINT ANTIBACTERIAL MOISTURIZING HAND SP - triclosan liquid
H E B

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 PERCENT

PURPOSE

ANTIBACTERIAL

USES

TO HELP REDUCE BACTERIA ON THE SKIN.

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 INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

APPLY TO WET HANDS, LATHER AND RINSE THOROUGHLY.

QUESTIONS OR COMMENTS

1-866-695-3030

INACTIVE INGREDIENTS

WATER, AMMONIUM LAURYL SULFATE, ACRYLATES COPOLYMER, COCAMIDOPROPYL BETAINE, GLYCOL DISTEARATE, GLYCERIN, SODIUM CHLORIDE, GLYCERETH-26, COCO-GLUCOSIDE, GLYCERYL OLEATE, LAURYL LACTYL LACTATE, GLYCERYL STEARATE, LAVENDULA ANGUSTIFOLIA (LAVENDER) FLOWER/LEAF/STEM EXTRACT, MENTHA PIPERITA (PEPPERMINT) LEAF EXTRACT, TOCOPHERYL ACETATE, BENZOPHENONE-4, DISODIUM EDTA, SODIUM HYDROXIDE, CITRIC ACID, BENZYL ALCOHOL, FRAGRANCE,

MANNITOL, CELLULOSE, HYDROXYPROPYL METHYLCELLULOSE, IRON OXIDES (CI 77491), EXT. VIOLET 2 (CI 60730), RED 33 (CI 17200), METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE.

WHITE LAVENDER AND MINT ANTIBACTERIAL MOISTURIZING HAND SP

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-292
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.3 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERETH-26 (UNII: NNE56F2N14)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL MONOOLEATE (UNII: 4PC054V79P)	
LAURYL LACTATE (UNII: G5SU0BFK7O)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
LAVANDULA ANGUSTIFOLIA FLOWERING TOP (UNII: 9YT4B71U8P)	
MENTHA PIPERITA LEAF (UNII: A389O33LX6)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
SULISOBENZONE (UNII: 1W6L629B4K)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:37808-292-08	236 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/30/2010	

Labeler - H E B (007924756)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture

Revised: 1/2011 H E B