

**WHITE LAVENDAR AND MINT ANTIBACTERIAL GENTLE FOAMING HAND SP -
triclosan liquid
HEB**

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.

DRUGS FACTS

PURPOSE

ANTIBACTERIAL

USES

FOR WASHING TO 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 ONTO DRY HANDS, WORK INTO RICH FOAMY LATHER, RINSE AND DRY THOROUGHLY.

QUESTIONS OR COMMENTS

1-866-695-3030

INACTIVE INGREDIENTS

WATER, SODIUM LAURETH SULFATE, DISODIUM LAURETH SULFOSUCCINATE, COCAMIDOPROPYLAMINE OXIDE, GLYCERIN, SODIUM CHLORIDE, PEG-7 GLYCERYL COCOATE, GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE, TETRASODIUM EDTA, PROPYLENE GLYCOL, BENZYL ALCOHOL, FRAGRANCE, TRIETHYLENE GLYCOL, PPG-1-PEG-9 LAURYL GLYCOL ETHER, LAVANDULA ANGUSTIFOLIA (LAVENDER) FLOWER/LEAF/STEM EXTRACT, MENTHA PIPERITA (PEPPERMINT) LEAF EXTRACT, ALOE BARBADENSIS LEAF JUICE, BENZOPHENONE-4, HYDROXYPROPYL METHYLCELLULOSE, EXT VIOLET 2 (CI 60730), RED 33 (CI 17200), METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE.

ACTIVE INGREDIENT

TRICLOSAN 0.3 PERCENT



orchid™

**WHITE
LAVENDER
& MINT**

**Antibacterial
Gentle Foaming
Hand Soap**

Moisturizing with Aloe Vera

Kills common germs

8 FLOZ (236 mL)

06-16723

Drug Facts

Active ingredient Purpose
Triclosan 0.3 % Antibacterial

Uses ■ For washing to 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 onto dry hands, work into rich foamy lather, rinse and dry thoroughly.

Questions/Comments?

1-866-695-3030

Inactive ingredients: Water (Aqua), Sodium Laureth Sulfate, Disodium Laureth Sulfosuccinate, Cocamidopropylamine Oxide, Glycerin, Sodium Chloride, PEG-7 Glycerol Cocoate, Guar Hydroxypropyltrimonium Chloride, Tetrasodium EDTA, Propylene Glycol, Benzyl Alcohol, Fragrance (Parfum), Triethylene Glycol, PPG-1-PEG-9 Lauryl Glycol Ether, Lavandula Angustifolia (Lavender) Flower/Leaf/Stem Extract, Mentha Piperita (Peppermint) Leaf Extract, Aloe Barbadosensis Leaf Juice, Benzophenone-4, Hydroxypropyl Methylcellulose, Ext Violet 2 (CI 60730), Red 33 (CI 17200), Methylchloroisothiazolinone, Methylisothiazolinone.

MADE IN CANADA

Distributed by: Parkway Manufacturing and Trading Company, San Antonio, TX 78218



0 41220 85877 4

06-16724

WHITE LAVENDAR AND MINT ANTIBACTERIAL GENTLE FOAMING HAND SP

triclosan liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-181
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)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DISODIUM LAURETH SULFO SUCCINATE (UNII: D6DH1DTN7E)	
CO CAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERYL COCOATE (UNII: WVK1CT5994)	
GUAR GUM (UNII: E89I1637KE)	
EDETATE SODIUM (UNII: MP1J8420LU)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
POLIDOCANOL (UNII: 0AWH8BFG9A)	
LAVANDULA ANGUSTIFOLIA FLOWERING TOP (UNII: 9YT4B71U8P)	
MENTHA PIPERITA LEAF (UNII: A389O33LX6)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SULISOBENZONE (UNII: 1W6L629B4K)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
METHYLCHLOROISO THIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISO THIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-181-08	236 mL in 1 BOTTLE		

Marketing Information

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

Labeler - HEB (007924756)**Registrant** - APOLLO HEALTH AND BEAUTY CARE (201901209)**Establishment**

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

Revised: 12/2010

HEB