

**CACTUS FLOWER AND MANGO ANTIBACTERIAL DEEP CLEANSING 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.

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.

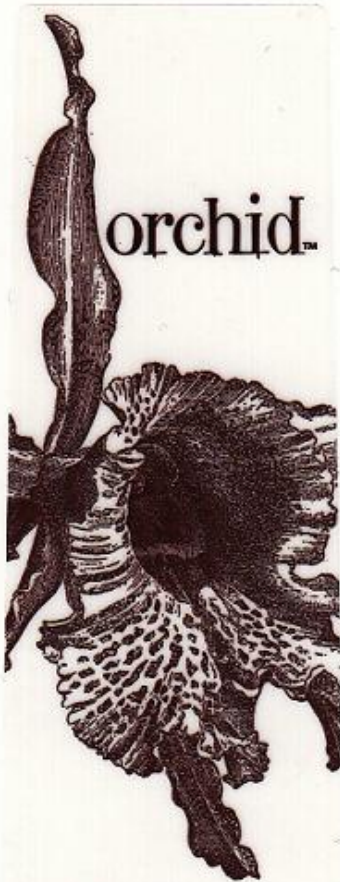
QUESTIONS OR COMMENTS

1-866-695-3030

INACTIVE INGREDIENTS

WATER, SODIUM LAURETH SULFATE, ACRYLATES COPOLYMER, COCAMIDOPROPYL BETAINE, COCAMIDE DEA, SODIUM CHLORIDE, FRAGRANCE, ALOE BARBADENSIS LEAF JUICE, PEG-7 GLYCERYL COCOATE, HELIANTHUS ANNUUS (SUNFLOWER) EXTRACT, CEREUS GRANDIFLORUS (CACTUS) FLOWER EXTRACT, MANGIFERA INDICA (MANGO) FRUIT EXTRACT, POLYETHYLENE, TETRASODIUM EDTA, SODIUM STYRENE/PEG-10 MALEATE/NONOXYNOL-10 MALEATE/ACRYLATES COPOLYMER, AMMONIUM

NONOXYNOL-4 SULFATE, CITRIC ACID, SODIUM HYDROXIDE, MANNITOL, CELLULOSE, HYDROXYPROPYL METHYLCELLULOSE, MICA, TITANIUM OXIDE (CI 77891), IRON OXIDES (CI 77491), IRON OXIDES (CI 77492), YELLOW 5 (CI 19140), RED 4 (CI 14700), RED 33 (CI 17200), METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE.



orchid™

CACTUS FLOWER & MANGO

Antibacterial Deep Cleansing Hand Soap with Exfoliating Beads Kills common germs

8 FLOZ (236 mL)

06-16721

Drug Facts

Active ingredient	Purpose
Triclosan 0.3 %	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/Comments?
1-866-695-3030

Inactive ingredients: Water (Aqua), Sodium Laureth Sulfate, Acrylates Copolymer, Cocamidopropyl Betaine, Cocamide DEA, Sodium Chloride, Fragrance (Parfum), Aloe Barbadosensis Leaf Juice, PEG-7 Glyceryl Cocoate, Helianthus Annuus (Sunflower) Extract, Cereus Grandiflorus (Cactus) Flower Extract, Mangifera Indica (Mango) Fruit Extract, Polyethylene, Tetrasodium EDTA, Sodium Styrene/PEG-10 Maleate/Nonoxynol-10 Maleate/Acrylates Copolymer, Ammonium Nonoxynol-4 Sulfate, Citric Acid, Sodium Hydroxide, Mannitol, Cellulose, Hydroxypropyl Methylcellulose, Mica, Titanium Oxide (CI 77891), Iron Oxides (CI 77491), Iron Oxides (CI 77492), Yellow 5 (CI 19140), Red 4 (CI 14700), Red 33 (CI 17200), Methylchloroisothiazolinone, Methylisothiazolinone.

MADE IN CANADA

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



CACTUS FLOWER AND MANGO ANTIBACTERIAL DEEP CLEANSING HAND SP

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-291
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: 059QF0K00R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
CO CAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERYL COCOATE (UNII: WVK1CT5994)	
HELIANTHUS ANNUUS FLOWER (UNII: BKJ0J3D1BP)	
SELENICEREUS GRANDIFLORUS FLOWER (UNII: I1877K4UNR)	
MANGO (UNII: I629IBNR86)	
LOW DENSITY POLYETHYLENE (UNII: J245LN42AD)	
EDETATE SODIUM (UNII: MP1J8420LU)	
NONOXYNOL-10 (UNII: K7O76887AP)	
AMMONIUM NONOXYNOL-4 SULFATE (UNII: 9HIA70O4J0)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MICA (UNII: V8A1AW0880)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-291-08	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