CACTUS FLOWER AND MANGO 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.

DRUG FACTS

ACTIVE INGREDIENT

TRICLOSAN 0.3 PERCENT

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, CEREUS GRANDIFLORUS (CACTUS) FLOWER EXTRACT,

MANGIFERA INDICA (MANGO) FRUIT EXTRACT, ALOE BARBADENSIS LEAF JUICE, BENZOPHENONE-4, HYDROXYPROPYL METHYLCELLULOSE, RED 4 (CI 14700), RED 33 (CI 17200), YELLOW 5 (CI 19140), METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE.

CACTUS FLOWER AND MANGO ANTIBACTERIAL GENTLE FOAMING HAND SP

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-182
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 SULFOSUCCINATE (UNII: D6DH1DTN7E)	
COCAMIDOPROPYLAMINE 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)	
SELENICEREUS GRANDIFLORUS FLOWER (UNII: II877K4UNR)	
MANGO (UNII: I629I3NR86)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SULISOBENZONE (UNII: 1W6L629B4K)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:37808-182-08	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		3019012093	manufacture

Revised: 12/2010 HEB