

**IMAGE ESSENTIALS ANTIBACTERIAL HAND MANDARIN AND GRAPEFRUIT-
triclosan liquid
K MART CORPORATION**

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 0.15%

PURPOSE

ANTIBACTERIAL

USES

FOR HAND WASHING TO DECREASE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER

STOP USING THIS PRODUCT AND ASK A DOCTOR IF

IRRITATION AND REDNESS DEVELOPS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY (1-800-222-1222)

DIRECTIONS

APPLY ONTO WET HANDS, WORK INTO A LATHER, RINSE THOROUGHLY

OTHER INFORMATION

STORE AT ROOM TEMPERATURE

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM LAURETH SULFATE, SODIUM CHLORIDE, DECYL GLUCOSIDE, COCAMIDOPROPYL BETAINE, COCAMIDE MEA, FRAGRANCE (PARFUM), PEG-120 METHYL GLUCOSE DIOLATE, PEG-18 GLYCERYL OLEATE/COCOATE, ALOE BARBADENSIS LEAF JUICE, POLYQUATERNIUM-7, DMDM HYDANTOIN, TETRASODIUM EDTA, CITRIC ACID, RED 33 (CI 17200), YELLOW 6 (CI 15985)

QUESTIONS OR COMMENTS?

1-800-842-7886

LABEL COPY



IMAGE ESSENTIALS ANTIBACTERIAL HAND MANDARIN AND GRAPEFRUIT

triclosan liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49738-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	1.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF301KX)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	

PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
PEG-18 GLYCERYL OLEATE/COCOATE (UNII: VD2D270332)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 160000 MW) (UNII: 0L414VCS5Y)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49738-015-08	222 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/23/2014	

Labeler - KMART CORPORATION (008965873)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(49738-015)

Revised: 11/2014

KMART CORPORATION