# IMAGE ESSENTIALS SPRING FRESH- triclos an liquid KMART 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.

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#### **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, FLUSH WITH WATER.

STOP USING THIS PRODUCT AND ASK A 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 (1-800-222-1222).

#### **DIRECTIONS**

APPLY ONTO WET HANDS. WORK INTO LATHER, RINSE THOROUGHLY AND DRY.

#### 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 DIOLEATE, PEG-18 GLYCERYL OLEATE/COCOATE, ALOE BARBADENSIS LEAF JUICE, POLYQUATERNIUM-7, DMDM HYDANTOIN, TETRASODIUM EDTA, CITRIC ACID, BLUE 1 (CI 42090), RED 33 (CI 17200).

#### **QUESTIONS OR COMMENTS?**

#### LABEL COPY





triclosan liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49738-298	
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: BPV390 UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6 Y)	
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0 K64F20 V)	
PEG-18 GLYCERYL OLEATE/COCOATE (UNII: VD2D270332)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
DMDM HYDANTO IN (UNII: BYR0546TOW)	
EDETATE SO DIUM (UNII: MP1J8420LU)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

P	ackaging			
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:49738-298-08	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	06/27/2013	

## 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(49738-298)

Revised: 7/2013 KMART CORPORATION