CVS PHARMACY ANTIBACTERIAL - triclos an soap CVS PHARMACY

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 USE AND ASK A DOCTOR

IF IRRITATION OR REDNESS DEVELOPS AND LASTS MORE THAN 7 DAYS.

KEEP OUT OF REACH OF CHILDREN

IF INGESTED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

DIRECTIONS

APPLY A SMALL AMOUNT ONTO WET HANDS. WORK INTO A RICH LATHER AND RINSE CLEAN.

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM C14-16 OLEFIN SULFONATE, LAURAMIDE DEA, SODIUM CHLORIDE, COCAMIDOPROPYL BETAINE, FRAGRANCE (PARFUM), CITRIC ACID, DMDM HYDANTOIN, GLYCERIN, TETRASODIUM EDTA, POLYQUATERNIUM-7, HYDROLYZED SILK PROTEIN, ALOE BARBADENSIS LEAF JUICE, RED 40 (CI 16035), YELLOW 5 (CI 19140), RED 33 (CI 17200).

LABEL COPY





CVS PHARMACY ANTIBACTERIAL

triclosan soap

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:59779-211

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X) TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9 W3D3YF5U)				
LAURIC DIETHANO LAMIDE (UNII: 129 12 VHG38)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)				
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)				
DMDM HYDANTOIN (UNII: BYR0546TOW)				
GLYCERIN (UNII: PDC6 A3C0 O X)				
EDETATE SO DIUM (UNII: MP1J8420LU)				
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600 KD) (UNII: 0L414VCS5Y)				
SILK, BASE HYDROLYZED (1000 MW) (UNII: UMQ31C11AY)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
FD&C YELLOW NO. 5 (UNII: 1753WB2F1M)				

D&C RED NO.33 (UNII: 9DBA0SBB0L)

I	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:59779-211-12	333 mL in 1 BOTTLE, PUMP				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	03/26/2012				

Labeler - CVS PHARMACY (062312574)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

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

Revised: 3/2012 CVS PHARMACY