

RITE AID RENEWAL ANTIBACTERIAL CLEAR SPRING- triclosan liquid
RITE AID 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 WASHING TO DECREASE BACTERIA ON THE SKIN.

WARNINGS

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT

STOP USING THIS PRODUCT AND ASK A DOCTOR IF

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

SQUEEZE ONTO A WET WASHCLOTH, SPONGE, POUF OR HANDS AND APPLY TO BODY. WORK INTO A RICH LATHER. RINSE THOROUGHLY.

QUESTIONS OR COMMENTS?

1-866-695-3030

INACTIVE INGREDIENTS

WATER (AQUA), SODIUM LAURETH SULFATE, COCAMIDOPROPYL BETAINE, PEG-8, FRAGRANCE (PARFUM), GLYCERIN, ISOSTEARAMIDOPROPYL MORPHOLINE LACTATE, POLYQUATERNIUM-10, COCAMIDOPROPYL PG-DIMONIUM CHLORIDE, DMDM HYDANTOIN, CITRIC ACID, TETRASODIUM EDTA, SODIUM CHLORIDE, BLUE 1 (CI 42090), RED 33 (CI 17200).



RITE AID RENEWAL ANTIBACTERIAL CLEAR SPRING			
triclosan liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-80 10
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)				
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
GLYCERIN (UNII: PDC6A3C0OX)				
ISOSTEARAMIDOPROPYL MORPHOLINE LACTATE (UNII: 082Y3WQI5K)				
POLYQUATERNIUM-10 (400 CPS AT 2 %) (UNII: HB1401PQFS)				
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE (UNII: 205Z54J075)				
DMDM HYDANTOIN (UNII: BYR0546TOW)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
EDETATE SODIUM (UNII: MP1J8420LU)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-8010-2	354 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	09/25/2012	

Labeler - RITE AID CORPORATION (014578892)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(11822-8010)

Revised: 9/2012

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