ORCHID - 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%

ANTIBACTERIAL

USES

TO HELP 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 TO WET HANDS, LATHER AND RINSE THOROUGHLY.

QUESTIONS/COMMENTS?

1-866-695-3030

INACTIVE INGREDIENTS:

WATER (AQUA), SODIUM LAURETH SULFATE, ACRYLATES COPOLYMER, GLYCOL DISTEARATE, MYRISTYL ALCOHOL, COCAMIDOPROPYL BETAINE, GLYCERIN, AMMONIUM CHLORIDE, GLYCERETH-26, COCO-GLUCOSIDE, GLYCERYL OLEATE, LAURYL LACTYL LACTATE, LAVANDULA ANGUSTIFOLIA (LAVENDER) FLOWER/LEAF/STEM EXTRACT, MENTHA PIPERITA (PEPPERMINT) LEAF EXTRACT, TOCOPHERYL ACETATE, BENZOPHENONE-4, DISODIUM EDTA, SODIUM HYDROXIDE, CITRIC ACID, BENZYL ALCOHOL, FRAGRANCE (PARFUM), MANNITOL, CELLULOSE, HYDROXYPROPYL METHYLCELLULOSE, IRON OXIDES (CI 77491), EXT. VIOLET 2 (CI 60730), RED 33 (CI 17200), MICA (CI 77019), TITANIUM DIOXIDE (CI 77891), METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE.

LABEL COPY

ORCHID

triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-297
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)	
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 1G56KV7BUJ)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMMONIUM CHLORIDE (UNII: 01Q9PC255D)	
GLYCERETH-26 (UNII: NNE56F2N14)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
LACTIC ACID (UNII: 33X04XA5AT)	
LAVANDULA ANGUSTIFOLIA FLOWERING TOP (UNII: 9YT4B71U8P)	
MENTHA PIPERITA LEAF (UNII: A389O33LX6)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SULISOBENZONE (UNII: 1W6L629B4K)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSE 2208 (4000 MPA.S) (UNII: 39J80LT57T)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
MICA (UNII: V8A1AW0880)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-297-08	236 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	04/12/2012	

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: 4/2012 HEB