CLOROX ANTIBACTERIAL HAND - WILD LAVENDER AND VANILLA-
chloroxylenol soap
Brand Buzz

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.

Clorox™ Antibacterial Hand Soap - Wild Lavender and Vanilla

DRUG FACTS

Active Ingredient

Chloroxylenol (0.3%)

Purpose

Antibacterial hand soap

Uses

- Helps to decrease bacteria on the skin.

Warnings

For external use only.

Keep out of reach of children, except under adult supervision. In case of accidental ingestion, drink a glass of water to dilute. If eye contact occurs, rinse thoroughly with water.

Directions

- Apply onto wet hands, lather for at least 30 seconds and rinse thoroughly.

Inactive Ingredients

WATER (AQUA, EAU), SODIUM LAURETH SULFATE, PEG-4 RAPESEEDAMIDE, PANTHENOL, TOCOPHERYL ACETATE, BUTYROSPERMUM PARKII (SHEA) BUTTER EXTRACT, COCOS NUCIFERA (COCONUT) FRUIT EXTRACT, COCAMIDOPROPYL BETAINE, PEG-150 DISTEARATE, POLYSORBATE 20, GLYCERIN, ASCORBIC ACID, BIOTIN, NIACINAMIDE, PROPYLENE GLYCOL, SODIUM CHLORIDE, CITRIC ACID, PEG/PPG-18/18 DIMETHICONE, SODIUM HYDROXIDE, TETRASODIUM EDTA, METHYLCHLOROISOTHIAZOLINONE, METHYLISOTHIAZOLINONE, FRAGRANCE (PARFUM), EXT. VIOLET 2 (CI 60730), RED 33 (CI 17200), ALPHA-ISOMETHYL IONONE, BUTYLPHENYL METHYLPROPIONAL

Questions?

1-855-539-3564

Distributed by Brand Buzz LLC,
1400 Broadway, New York, NY 10018

PRINCIPAL DISPLAY PANEL - 295 mL Bottle Label

***CLOROX*™**

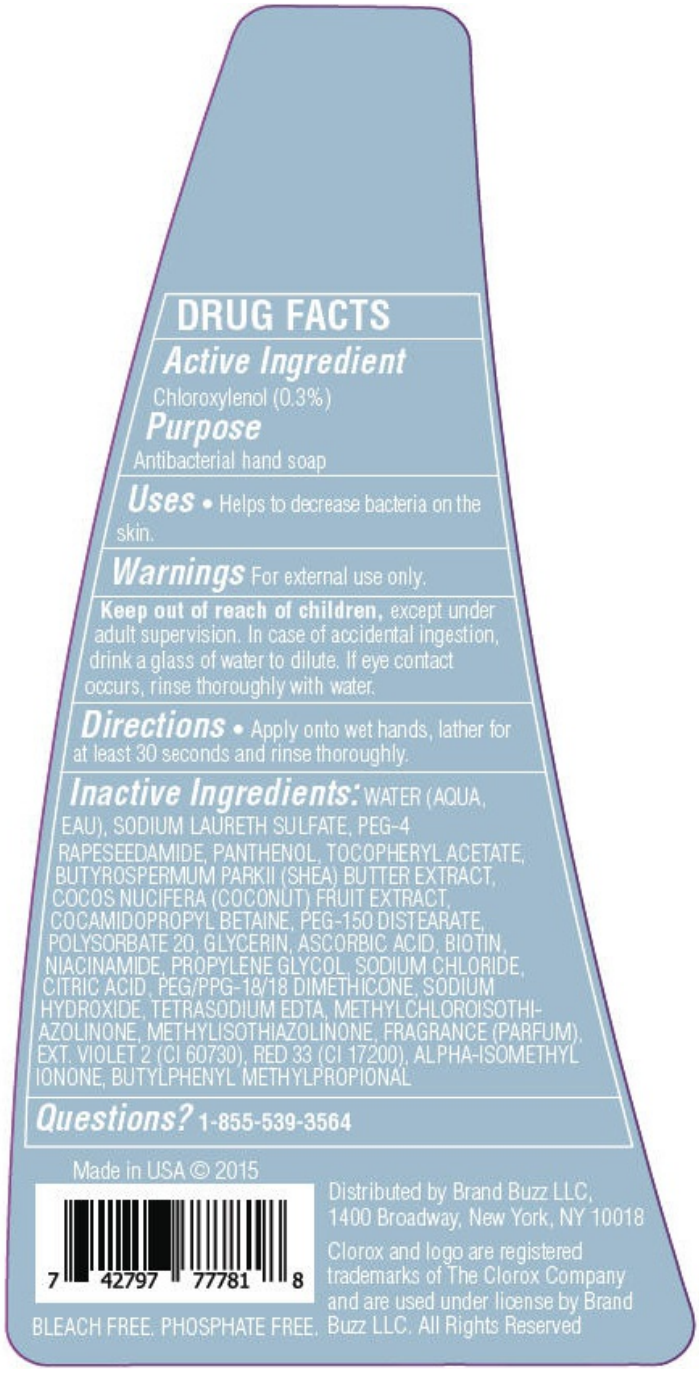
antibacterial

HAND SOAP
DERMATOLOGIST TESTED

MOISTURIZING

wild lavender
& vanilla

10 FL OZ (295 mL)



CLOROX ANTIBACTERIAL HAND - WILD LAVENDER AND VANILLA

chloroxylenol soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69540-0075
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Chloroxylenol (UNII: 0F32U78V2Q) (Chloroxylenol - UNII:0F32U78V2Q)	Chloroxylenol	0.003 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Sodium Laureth-3 Sulfate (UNII: BPV390UAP0)	
PEG-4 Rapeseedamide (UNII: 89575CN928)	
Panthenol (UNII: WV9CM0O67Z)	
.Alpha.-Tocopherol Acetate (UNII: 9E8X80D2L0)	
Sheanut Oil (UNII: O88E196QRF)	
Coconut (UNII: 3RT3536DHY)	
Cocamidopropyl betaine (UNII: 5OCF3O11KX)	
PEG-150 Distearate (UNII: 6F36Q0I0AC)	
Polysorbate 20 (UNII: 7T1F30V5YH)	
Glycerin (UNII: PDC6A3C0OX)	
Ascorbic Acid (UNII: PQ6CK8PD0R)	
Biotin (UNII: 6SO6U10H04)	
Niacinamide (UNII: 25X51I8RD4)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Sodium Chloride (UNII: 451W47IQ8X)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
PEG/PPG-18/18 Dimethicone (UNII: 9H0AO7T794)	
Sodium Hydroxide (UNII: 55X04QC32I)	
Edetate Sodium (UNII: MP1J8420LU)	
Methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
Methylisothiazolinone (UNII: 229D0E1QFA)	
EXT. D&C Violet NO. 2 (UNII: G5UX3K0728)	
D&C Red NO. 33 (UNII: 9DBA0SBB0L)	
Isomethyl-.Alpha.-Ionone (UNII: 9XP4LC555B)	
Butylphenyl Methylpropional (UNII: T7540GJV69)	

Product Characteristics

Color	PURPLE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69540-0075-1	295.735 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	02/01/2016	

Labeler - Brand Buzz (079266204)

Revised: 1/2016 Brand Buzz