CLOROX ANTIMICROBIAL- chloroxylenol soap The Clorox Company

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.

Active Ingredient

Chloroxylenol 0.5% w/w

Purpose

Antimicrobial

Uses

- For handwashing to decrease bacteria on the skin
- Ideal for repeated washing

Warnings

For external use only

When using this product

- **do not use** in or near eyes
- discontinue use if irritation and redness develop, or if condition persists for more than 72 hours

KEEP OUT OF REACH OF CHILDREN. In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

Directions

- Wet hands
- Apply product
- Wash hands thoroughly
- Rinse vigorously

Other Information

Store at room temperature

*Inactive Ingredients*Cocoamidopropyl Betaine, DMDM Hydantoin, Ethylene Glycol Monostearate, FD&C Blue #1, Fragrance, Methylchloroisothiazolinone, Methylisothiazolinone, Methylparaben, PEG-120 Methyl Glucose Dioleate, Propylparaben, Sodium Chloride, Sodium Laureth Sulfate, Water.

Questions or Comments?

QUESTIONS OR COMMENTS?

Call 1-800-638-2625 or visit www.hlk.cc

Clorox Antimicrobial Hand Soap

18 oz (532 mL)

Drug Facts (Continued)

Directions ■ Wet hands
■ Wash hands thoroughly
■ Rinse vigorously

Other information

Inactive ingredients Cocoamidopropyl Betaine, DMDM Hydantoin, Ethylene Glycol Monostearate, FD&C Blue #1, Fragrance, Methylchloroisothiazolinone, Methylisothiazolinone, Methylparaben, PEG-120 Methyl Glucose Dioleate, Propylparaben, Sodium Chloride, Sodium Laureth Sulfate, Water.

QUESTIONS OR COMMENTS? Call 1-800-638-2625 or

visit www.hlk.cc.
For more product ingredient information, visit www.lngredientsinside.com.
For SDS, go to: www.cloroxprofessional.com.
Made in the U.S.A.

Mfd. for Clorox Professional Products Company, 1221 Broadway, Oakland, CA 94612



CLOROX ANTIMICROBIAL

chloroxylenol soap

Prod	nct	Info	rma	tion

Product Type HUMAN OTC DRUG Item Code (Source) NDC:26509-0011

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)	
METHYLPARABEN (UNII: A218 C7H19 T)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
DMDM HYDANTO IN (UNII: BYR0546 TOW)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
METHYLCHLORO ISO THIAZO LINO NE (UNII: DEL7T5QRPN)	
WATER (UNII: 059QF0KO0R)	
METHYLISO THIAZO LINO NE (UNII: 229 D0 E1QFA)	

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:26509-0011-5	$532\ mL$ in $1\ BOTTLE;$ Type $0\colon Not\ a\ Combination\ Product$	0 1/25/20 16			
Marketing Inf	ormation				
Marketing Inf		Marketing Start Date	Marketing End Date		
	ry Application Number or Monograph Citation	Marketing Start Date 0 1/25/20 16	Marketing End Date		

Labeler - The Clorox Company (009138033)

Registrant - Carroll Company (007372329)

Establishment				
Name	Address	ID/FEI	Business Operations	
Carroll Co.		007372329	manufacture (26509-0011)	

Revised: 1/2016 The Clorox Company