CHLOROXYLENOL- chloroxylenol liquid
Vi-Jon, LLC

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.

Mountain Falls 710.002/710AC rev 1 Citrus Orange Antibacterial Hand Soap

Active ingredient

Chloroxylenol 0.3%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hands only

When using this product

- do not get into eyes. If contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Conrol Center right away.

Directions

- wet hands
- apply palmful to hands,
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

water, sodium C14-16 olefin sulfonate, lauramine oxide, sodium laureth sulfate, sodium lauryl sulfate, isopropylideneglycerol, sodium chloride, alcohol denat., phenoxyethanol, citric acid, methylisothiazolinone, trisodium ethylenediamine disuccinate, fragrance, yellow 5, red 33

Do not add bleach. Not for use in dishwashers.

Contains Surfactants. Phosphate Free

disclaimer

This product is not manufactured or distributed by Procter & Gamble, distributor of Dawn Ultra Antibacteial Hand Soap Orange Scent

principal display panel

mountain

falls

*Compare

to Dawn

tough

on

grease

ultra

concentrated

cleaning

power

dish soap

antibacterial hand soap

orange scent

24 FL OZ (710 mL)

CHLOROXYLENOL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-710
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3.09 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SOLKETAL (UNII: 3XK098O8ZW)	
sodium chloride (UNII: 451W47IQ8X)	
ALCOHOL (UNII: 3K9958V90M)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
trisodium ethylenediamine disuccinate (UNII: YA22H34H9Q)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-710-50	710 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/14/2017	
2	NDC:11344-710-57	1183 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/14/2017	
3	NDC:11344-710-51	1770 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/14/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/14/2017	

Labeler - Vi-Jon, LLC (790752542)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(11344-710)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11344-710)

Revised: 4/2022

Vi-Jon, LLC