CHLOROXYLENOL- chloroxylenol liquid
Old East Main Co

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.

TrueLiving 560.002/560AC-AD-AE 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 Control Center right away.

Directions

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

Inactive ingredients

water, sodium C14-16 olefin sulfonate, lauramine oxide and/or lauramidopropylamine oxide and myristamidopropylamine oxide, sodium laureth sulfate, sodium lauryl sulfate, sodium chloride, sodium xylenesulfonate, alcohol denat., fragrance, limonene, phenoxyethanol, citric acid, methylisothiazolinone, tetrasodium EDTA, yellow 5, red 33

Do not add bleach.

Not for use in dishwashers.

Contains Surfactants Phosphate Free

**When compared to non-concentrated formulas

*This product is not manufactured or distributed by Procter & Gamb;le, distributor of Dawn Ultra Antibacterial Hand Soap Orange Scent

principal display panel

ULTRA CONCENTRATED 3X CLEANING POWER

trueliving

ULTRA

Antibacterial

Hand Soap

Dishwashing Liquid

TOUGH ON GREASE

Compare to Dawn Ultra Antibacterial Hand Soap Orange Scent

CITRUS ORANGE SCENT

18 FL OZ (532 mL)

CHLOROXYLENOL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-984
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)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALCOHOL (UNII: 3K9958V90M)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
DITETRACYCLINE TETRASODIUM EDETATE (UNII: WX0A0IT7K5)	
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:55910-984-44	532 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/01/2020	
2	NDC:55910-984-57	1183 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/01/2020	

Labeler - Old East Main Co (068331990)

Registrant - Vi-Jon,LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(55910-984)

Revised: 3/2022 Old East Main Co