INODERM CHOICE ANTIBACTERIAL- chloroxylenol liquid
Avro Enterprises LLC

Inoderm Choice Antibacterial Soap

Drug Facts

Active ingredient

Chloroxylenol 0.3% w/w

Purpose

Antiseptic

Uses

- Handwash to decrease bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

Warnings

For external use only

When using this product

keep away from eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if

irritation or redness develop or if 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 with water and dispense small amount of product into cupped palm of hand.
- Lather vigorously for at least 15 seconds.
- Rinse with water and dry thoroughly.

Inactive ingredients

Water, Sodium Lauryl Ether Sulfate, Sodium Chloride, Cocamide MIPA, Sodium Sulfate, Fragrance, Magnesium Nitrate, Citric Acid, FD&C Yellow # 5, Methylchloroisothiazolinone, Magnesium Chloride, Methylsothiazolinone, FD&C Red # 4

Package Labeling:

INODERM CHOICE ANTIBACTERIAL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73062-523
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	

METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73062-523-42	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/14/2023	

Labeler - Avro Enterprises LLC (804030166)

Revised: 10/2023 Avro Enterprises LLC