MICRO-ASEPTIC- chloroxylenol liquid
ASEPTIC CONTROL PRODUCTS, INC.

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.

Micro-Aseptic 81591-001

Active Ingredient

PCMX (para-chloro-meta-xylenol) 0.6%

Purpose

Antimicrobial

Use

for hand-washing to decrease bacteria on the skin

Warnings

For external use only

When using this product
- do not get into eyes
- of contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if
- irritation and redness develop

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control center right away

Directions
- ready to use
- do not dilute
- wet hands and forearms
- apply 5 milliters (teaspoonful) or palmful to hands and forearms
- scrub thoroughly for 1 minute and rinse

Inactive Ingredients

Water, Sodium Lauryl Sulfate, Sodium Laureth Sulfate, Cocamidopropylbetaine, Propylene Glycol, Tetrasodium EDTA, Isopropanol, Lauramine oxide, Boric Acid, PEG-75

Lanolin, DMDM Hydantoin, Fragrance, Aloe Vera, FD&C Blue #1, FD&C Yellow #5

Micro-

Aseptic

Antimicrobial

Skin Cleanser

NDC # 81591-001-02

Manufactured for

ASEPTIC CONTROL PRODUCTS, INC

1225 Carnegie Street, Unit 104

Rolling Meadows, IL 60008

Made in the U.S.A.

Item #: 1955-16OZ

473 ml (16 FL. OZ.)

Micro-Aseptic

Antimicrobial Skin Cleanser

NDC # 81591-001-02

Manufactured for

ASEPTIC CONTROL PRODUCTS, INC

1225 Carnegie Street, Unit 104
Rolling Meadows, IL 60008
(800) 448-0131
Fax: (847) 342-1809

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Drug Facts

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MICRO-ASEPTIC

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81591-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.6 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
EDETATE SODIUM (UNII: MP1J8420LU)	
BORIC ACID (UNII: R57ZHV85D4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81591-001-01	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2021	
2	NDC:81591-001-02	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2021	
3	NDC:81591-001-03	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/26/2021	

Labeler - ASEPTIC CONTROL PRODUCTS, INC. (161977020)

Establishment

Name	Address	ID/FEI	Business Operations
Enzyme Solutions, Inc.		004994559	manufacture(81591-001)