

PURO ANTIBACTERIAL WASH FOAM- chloroxylenol solution
CWGC LA 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.

Puro Antibacterial Wash Foam

Active ingredient

Chloroxylenol, 0.5%

Purpose

Antibacterial

Uses

- For handwashing to reduce bacteria on the skin

Warnings

For external use only

When using this product

- Avoid contact with eyes. In case of eye contact, flush with water

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply foaming cleanser to dry hands
- Rub hands together to spread lather
- Wash for 15-20 seconds
- Rinse & dry hands thoroughly

Inactive ingredients

Water, Sodium Laureth Sulfate, Isopropyl Alcohol, Cocamidopropyl Betaine, Decyl Glucoside, SODIUM C14-16 OLEFIN SULFONATE, Hydrogenated castor Oil, DMDM Hydantoin, Citric Acid, Fragrance, FD&C Blue No.1, FD&C Yellow No.5.



FOAM Antibacterial WASH

Drug Facts

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Sierra Health, Inc. Placentia, CA 92870 855.686.1888



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Item # HS871161

1L 2
33.8 fl. oz.

ANTIBACTERIAL WASH

PURO ANTIBACTERIAL WASH FOAM

chloroxylenol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70415-401
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70415-401-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/04/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/04/2016	

Labeler - CWGC LA Inc. (034967904)

Revised: 12/2018

CWGC LA Inc.