CLEANSLATE ANTIBACTERIAL FOAM- chloroxylenol liquid
Cleanslate Group 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.

Cleanslate Foam Hand 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, rinse eyes with water.

Stop use and ask a doctor if irritation or redness develop or last 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 sufficient amount of product into cupped palm of hand
- wash thoroughly for 15 seconds
- rinse under running water and dry thoroughly

Inactive ingredients

water, sodium laureth sulfate, cocamide DEA, DMDM hydantoin, ethyl alcohol, phenoxyethanol, isopropyl alcohol, fragrance, citric acid, FD&C red 4

MFD FOR CLEANSLATE GROUP LLC

1420 EAST LINDEN AVE

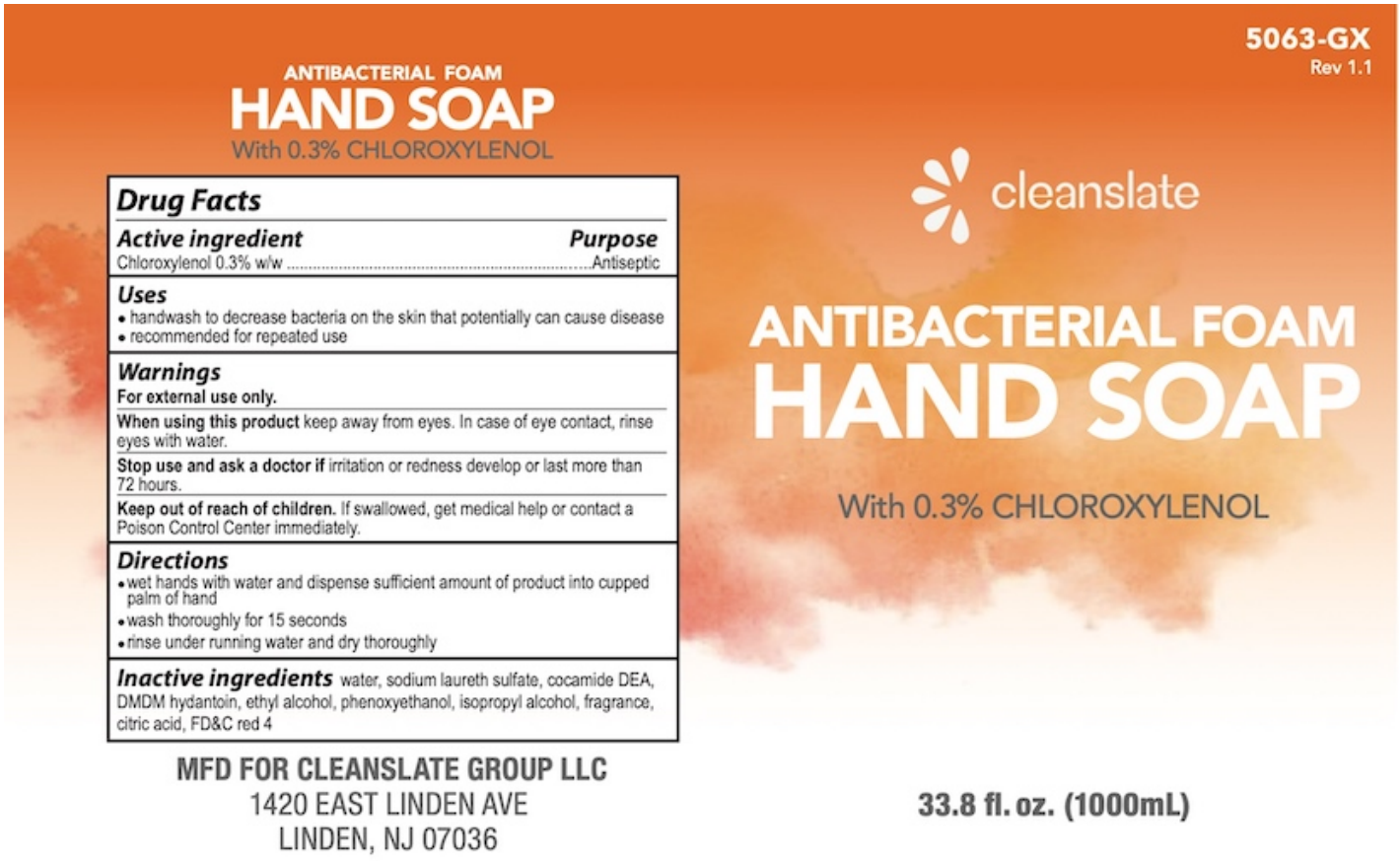
LINDEN, NJ 07036

cleanslate

ANTIBACTERIAL FOAM HAND SOAP

With 0.3% CHLOROXYLENOL

33.8 fl. oz. (1000mL)



CLEANSLATE ANTIBACTERIAL FOAM

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80586-523
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.3 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALCOHOL (UNII: 3K9958V90M)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80586-523-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	11/17/2021	07/26/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/17/2021	07/26/2024

Labeler - Cleanslate Group LLC (117657934)

Revised: 1/2023 Cleanslate Group LLC