SOFTEN SURE FOAM ANTIMICROBIAL- chloroxylenol liquid
Best Sanitizers, 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.

Soften Sure Foam Soap

Active Ingredient

Chloroxylenol 1.0%

Purpose

Antimicrobial Hand Cleanser

Uses

- Handwash to help reduce bacteria that potentially can cause diseases
- Helps to prevent cross contaminiation by hand contact
- Helps to prevent drying of the skin
- Recommended for repeated use

Warnings

- **For external use only**

When Using this product

- Do not use in or near eyes.

Keep out of reach of children

In case of accidental ingestion seek medical attention or contact a poison center immediately.

Directions

- Wet hands and forearms
- Apply an application to hands and forearms
- Scrub thoroughly for at least 30 seconds (include the fingernails and the cuticles)
- Rinse

Other Information

- Store in a cool dry place below 104° F

Inactive ingredients

purified water, potassium cocoate, tetrasodium EDTA, FDC No. 1, Yellow No.5, fragrance

Questions?

Contact **Best Sanitizers** Mon-Fri 9am - 4pm PST at **888-225-3267**

NDC 59900-211-06

NDC 59900-211-10

NDC 59900-211-11

Soften Sure Foam Soap antimicrobial

SOFTEN SURE FOAM ANTIMICROBIAL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59900-211
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	10 g in 1000 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	
EDETATE SODIUM (UNII: MP1J8420LU)	

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:59900-211-06	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2008	
2	NDC:59900-211-10	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2008	
3	NDC:59900-211-11	1000 mL in 1 BAG; Type 0: Not a Combination Product	09/01/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/01/2008	

Labeler - Best Sanitizers, Inc (957473614)

Establishment

Name	Address	ID/FEI	Business Operations
Best Sanitizers, Inc		627278224	manufacture(59900-211)

Revised: 1/2019 Best Sanitizers, Inc