PRESTIGE- chloroxylenol soap
Sunburst Chemicals, Inc.

Prestige

Active Ingredient

Chloroxylenol 1% w/w

Purpose

Skin Antimicrobial

Use

reduces amount of bacteria on hands

Warnings

For external use only. Do not use in eyes.

Discontinue use if irritation and redness develop. If condition persists for more than 72 hours, consult a physician.

Not for use on children under 6 months of age.

For institutional and professional use only.

Directions

- Wet hands and forearms.
- Apply a small amount (5 mL) or palmful to hands and forearms.
- Scrub thoroughly for at least fifteen seconds.
- Rinse completely and dry.

Inactive Ingredients

Water, Sodium C14-16 Olefin Sulfonate, Sodium Laureth Sulfate, Lauramide DEA, Isopropyl Alcohol, Polyquaternium-7, Fragrance, Styrene/Acrylates Copolymer, Tetrasodium EDTA, Citric Acid

Prestige is a skin care hand wash for healthcare, food service, and institutional personnel to help reduce the amount of bacteria and fungus on hands. A high quality hand soap that is a gentle cleaner for frequent use.

Drug Facts

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Please Recycle the Container

LBL1327-2.3

Manufactured For:
SUNBURST CHEMICALS, INC.
Minneapolis, MN 55420
(952) 884-3144
www.sunburstresults.com

PRESTIGE

chloroxylenol soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63621-335
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	

Product Characteristics			
Color	white (white, opaque liquid)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63621-335-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/03/2020	

Labeler - Sunburst Chemicals, Inc. (006159339)

Revised: 3/2024 Sunburst Chemicals, Inc.