SUNDROPS 77- benzalkonium chloride soap
Sunburst Chemicals, Inc.

SunDrops 77

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

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

Directions

- Pump a small amount of foam into palm of hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, Glycerin, Propylene Glycol, Phenoxyethanol, Lauramine Oxide, Panthenol, Fragrance, Citric Acid

SUNDROPS 77

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63621-354
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PANTHENOL (UNII: WV9CM0O67Z)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Product Characteristics

Color	white (water white - colorless, dispensed as a white foam)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63621-354-85	8 in 1 BOX	09/28/2012	
1		1000 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:63621-354-70	6 in 1 BOX	05/08/2008	06/30/2022
2		500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/05/2008	

Labeler - Sunburst Chemicals, Inc. (006159339)

Revised: 10/2024 Sunburst Chemicals, Inc.