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, Cocamidopropyl PG-Dimonium Chloride Phosphate, Dihydroxyethyl Cocamine Oxide, Acetamidoethoxyethanol, 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
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
ACETAMIDOETHOXYETHANOL (UNII: LVX2APC4XR)	
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: 5/2024 Sunburst Chemicals, Inc.