SUNDROPS 87- benzalkonium chloride soap
Sunburst Chemicals, Inc.

SunDrops 87

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Skin Antimicrobial

Use

Reduces amount of bacteria on hands

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.
- Scrub thoroughly for at least fifteen seconds.
- Rinse completely and dry.

Inactive Ingredients

Water, Lauramine Oxide, Glycerin, PEG-120 Methyl Glycose Dioleate, DMDM Hydantoin, Citric Acid, Fragrance

SUNBURST™

SunDrops 87

Foaming Antibacterial Hand Soap

Distributed Exclusively By:
Sunburst Chemicals, Inc.
Minneapolis, MN 55420
www.sunburstresults.com

Please Recycle This Container

Contains: Eight Bags

Net Contents Each: 33.8 fl. oz. (1 qt. 1.8 fl. oz.) 1000 mL
Total Net Volume: 270.4 fl. oz. (2 gal. 14.4 fl. oz.) 8 L

LBL1714-3.1

Drug Facts

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LOT #:
EXP:

Stock Number
3538708

10784331199400

SUNDROPS 87

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63621-387
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**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Product Characteristics			
Color	white (colorless - water-white, 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:63621-387-50	6 in 1 BOX	06/08/2020	
1		500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		
2	NDC:63621-387-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020	11/17/2023
3	NDC:63621-387-65	8 in 1 BOX	07/14/2020	
3		1000 mL in 1 BAG; 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)	06/08/2020	

Labeler - Sunburst Chemicals, Inc. (006159339)

Revised: 1/2025 Sunburst Chemicals, Inc.