

**SUNDROPS 87- benzalkonium chloride soap**  
**Sunburst Chemicals, Inc.**

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**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.

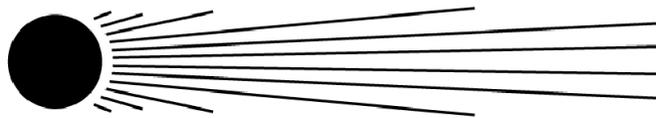
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

**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

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



LOT #: EXP:

Stock Number 3538708



LBL1714-3.1

10784331199400

<b>Drug Facts</b>	
<b>Active Ingredient</b>	<b>Purpose</b>
Benzalkonium Chloride 0.1% .....	Skin Antimicrobial
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Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.	
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<b>Directions</b>	
<ul style="list-style-type: none"> <li>• Pump a small amount of foam into palm of hand.</li> <li>• Scrub thoroughly for at least fifteen seconds.</li> <li>• Rinse completely and dry.</li> </ul>	
<b>Inactive Ingredients</b>	
Water, Lauramine Oxide, Glycerin, PEG-120 Methyl Glucose Dioleate, DMDM Hydantoin, Citric Acid, Fragrance	

**SUNDROPS 87**

benzalkonium chloride soap

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63621-387
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

Ingredient Name	Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>LAURAMINE OXIDE</b> (UNII: 4F6FC4MI8W)	
<b>PEG-120 METHYL GLUCOSE DIOLEATE</b> (UNII: YM0K64F20V)	
<b>DMDM HYDANTOIN</b> (UNII: BYR0546TOW)	

## Product Characteristics

<b>Color</b>	white (colorless - water-white, dispensed as a white foam)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## 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.