

**SUNDROPS 77- benzalkonium chloride soap**  
**Sunburst Chemicals, Inc.**

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**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



**Foaming Instant Hand Sanitizer  
ALCOHOL FREE**

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



LOT #:  
EXP:

**Stock Number**  
3537750



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

Drug Facts	
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## SUNDROPS 77

benzalkonium chloride soap

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63621-354
<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>PANTHENOL</b> (UNII: W9CM0067Z)	
<b>PHENOXYETHANOL</b> (UNII: H1E492ZZ3T)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>LAURAMINE OXIDE</b> (UNII: 4F6FC4MI8W)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

### Product Characteristics

<b>Color</b>	white (water white - colorless, 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:63621-354-85	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		
3	NDC:63621-354-40	4 in 1 BOX	10/16/2025	
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)	05/05/2008	

**Labeler** - Sunburst Chemicals, Inc. (006159339)

Revised: 11/2025

Sunburst Chemicals, Inc.