SUNDROPS 75- alcohol gel
Sunburst Chemicals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

SunDrops 75

Active Ingredient

Ethyl Alcohol 60% v/v

Purpose

Skin Sanitizer

Use

To help reduce amount of bacteria on the skin

Warnings

Flammable. Keep away from fire or flame.

For external use only. If swallowed, seek medical attention.

When using this product do not use around or near the eyes. If contact occurs, flush eyes with water and contact doctor immediately.

Stop use and consult a doctor when skin irritation appears and lasts.

Keep out of reach of children.

Directions

- Place 5 grams or palmful of product in one hand.
- Spread on hands and rub into skin until dry.
- Place a smaller amount (2.5 grams) in one hand & spread over hands & wrists.
- Rub into skin until dry.
- Supervise children while using this product.

Inactive Ingredients

Water, Glycerin, Fragrance, Carbomer, Propylene Glycol, Diisopropylamine

SUNBURST CHEMICALS

SunDrops 75 - 8 fl. oz.

SUNDROPS 75

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63621-356
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	0.6 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

Product Characteristics

Color	white (water white - colorless, crystal clear)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63621-356-20	8 in 1 BOX	10/14/2009	
1		1000 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:63621-356-25	12 in 1 BOX	10/14/2009	
2		236.6 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination		

2		Product		
3	NDC:63621-356-30	6 in 1 BOX	07/20/2016	
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 not final	part333E	10/14/2009	

Labeler - Sunburst Chemicals, Inc. (006159339)

Revised: 7/2016 Sunburst Chemicals, Inc.