HAND SANITIZER- alcohol aerosol, spray Cosmetic Specialty Labs, 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.

Active Ingredient

SD Alcohol 40 75%

Antimicrobial

Use

Hand Sanitizer to help reduce bacteria on the skin.

Warnings

Flammable. Keep away from fire or flame. For external use only.

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughtly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

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

Directions

- Use enough product in your palm to cover hands and rub hands together biskly until dry.
- Children under 6 years of age should be supervised when using this product.

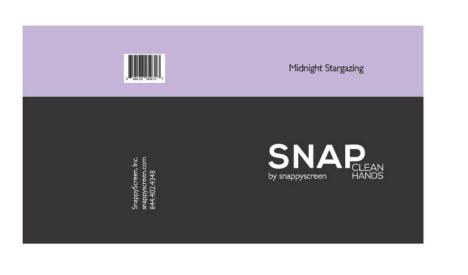
Other information

- Store below 110F (43C)
- May discolor fabrics or surfaces

Inactive ingredients

Aloe Barbadensis Leaf Juice, Caprylyl Glycol, Fragrance, Glycerin, Phenoxyethanol, Polysorbate-20, Purified Water, Tocopheryl Acetate.

Package Label Outer box







Package Label Refill bottle





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HAND SANITIZER							
alcohol aerosol, spray							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:58133-959			
Route of Administration	TOPICAL						
Active Ingredient/Active	Molety						
Ingredie	nt Name		Basis of Strength	Strength			
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	75 mL in 100 mL			
lus stins lusure dis sta							
Inactive Ingredients							
	Strength						
POLYSORBATE 20 (UNII: 7T1F30V							
PHENOXYETHANOL (UNII: HIE492ZZ3T)							
WATER (UNII: 059QF0K00R)							
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)							
CAPRYLYL GLYCOL (UNII: 00YIU5438U)							
GLYCERIN (UNII: PDC6A3C0OX)							
ALOE VERA LEAF (UNII: ZY81Z83H0X)							

Packaging							
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:58133-959- 72	1 in 1 BOX	05/03/2023				
1	NDC:58133-959- 69	178 mL in 1 BOTTLE; Type 0: Not a Combination Product					
Marketing Information							
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OT fin	C monograph no al	t part333A	05/03/2023				

Establishment								
Name	Address	ID/FEI	Business Operations					
Cosmetic Specialty Labs, Inc.		032973000	manufacture(58133-959)					

Revised: 5/2023

Cosmetic Specialty Labs, Inc.