BSAFE3 HAND SANITIZER- alcohol gel Active Creations LLC

BSAFE3 Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only-hands

Flammable. Keep away from heat and flame

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact Poison Control Center right away

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use under adult supervision
- not recommended for infants

Other Information

do not store above 105°F

- may discolor some fabrics
- harmful to wood finishes and plastics

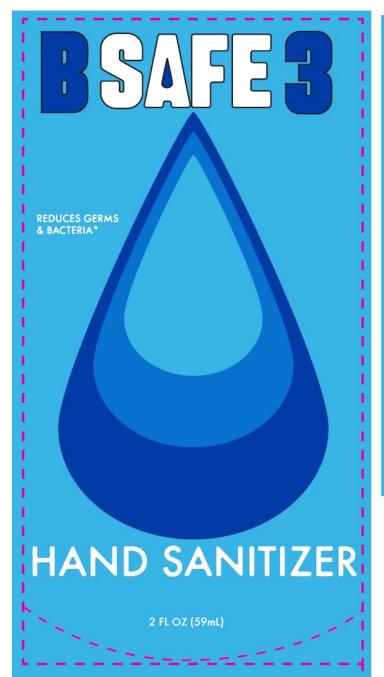
Inactive ingredients

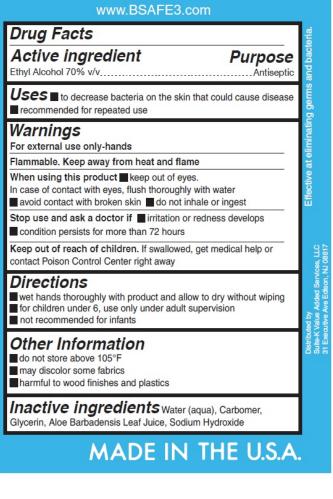
Water (aqua), Carbomer, Glycerin, Aloe Barbadensis Leaf Juice, Sodium Hydroxide

Package Labeling: 68957-2000-0



Package Labeling:68957-2000-1

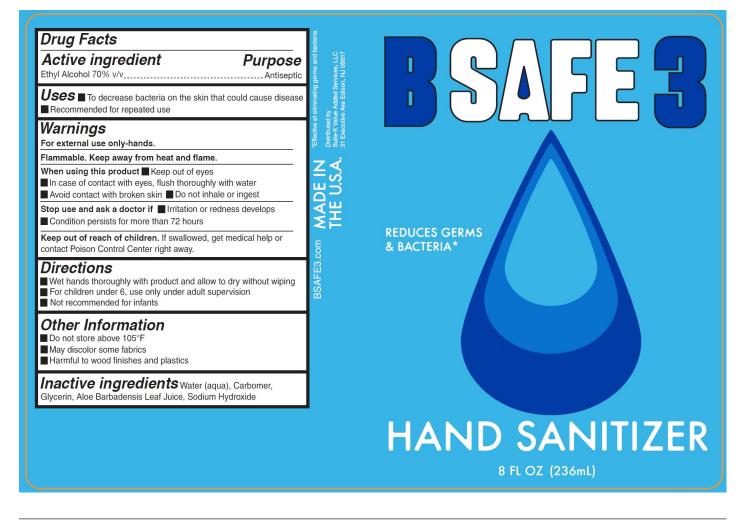




Package Labeling:68957-2000-2



Package Labeling:68957-2000-3



BSAFE3 HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68957-2000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
GLYCERIN (UNII: PDC6A3C0OX)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68957- 2000-0	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:68957- 2000-1	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
3	NDC:68957- 2000-2	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
4	NDC:68957- 2000-3	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/01/2020	

Labeler - Active Creations LLC (119119545)

Establishment				
Name	Address	ID/FEI	Business Operations	
Active Creations LLC		119119545	manufacture(68957-2000)	

Revised: 1/2024 Active Creations LLC