

CAREONE INSTANT HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E- ethyl alcohol liquid

American Sales Company

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.

Drug Facts

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- to help reduce bacteria on the skin.

Warnings

For external use only. Flammable. Keep away from source of heat or fire.

When using this product

- avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

- irritation or redness develops and lasts.

Keep out of reach of children.

- In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

- put enough product in your palm to cover hands and rub hands together until dry.
- children under 6 years should be supervised when using this product.

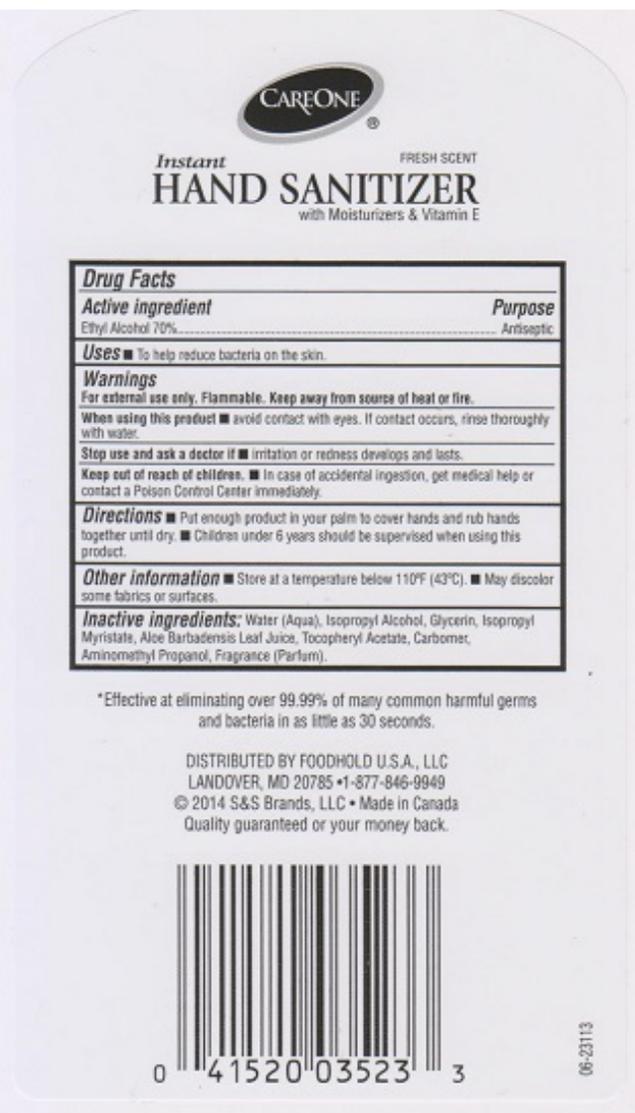
Other information

- store at a temperature below 110°F (43°C)
- may discolor some fabrics or surfaces

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Glycerin, Isopropyl Myristate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate, Carbomer, Aminomethyl Propanol, Fragrance (Parfum).

Label Copy



CAREONE INSTANT HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41520-178
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
CARBOMER 934 (UNII: Z135WT9208)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41520-178-16	443 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/25/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/25/2017	

Labeler - American Sales Company (809183973)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(41520-178)

Revised: 4/2017

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