

CAREONE VITAMIN E HAND SANITIZER- 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 65%

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 certain 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



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06-21876



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06-21882

CAREONE VITAMIN E HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41520-059	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	650 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0K00R)				
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:41520-059-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/25/2017	
2	NDC:41520-059-02	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/25/2017	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	09/25/2017		

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(41520-059)