# CAREONE VITAMIN E HAND SANITIZER- ethyl alcohol gel Retail Business Services, LLC

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.

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### **Drug Facts**

### **Active Ingredients**

Ethyl Alcohol (65%)

### **Purpose**

Antiseptic

#### Uses

For personal hand hygiene to help prevent the spread of bacterial, and can be used in place of hand washing if soap and water are not available.

## 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. Do not inhale.

#### Do not use

- on children less than 2 months of age
- on open skin wounds
- on broken or damaged skin
- if you are pregnant or breastfeeding

#### 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**

- For occasional and personal use
- Rub thoroughly into hands for at least 30 seconds. Allow to 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, PEG-6, Fragrance, Isopropyl Alcohol, Glycerin, Isopropyl Myristate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate, AMP-Acrylates/Vinyl Isodecanoate Crosspolymer., Yellow 5 (Cl 19140), Blue 1 (Cl 42090)

59 ml

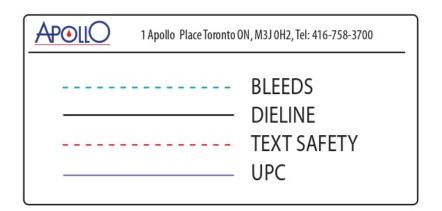
# Die 1.56 in x 2.79 in



Front Label



**Back Label** 

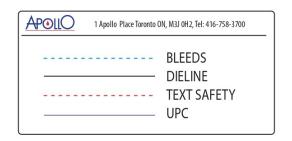


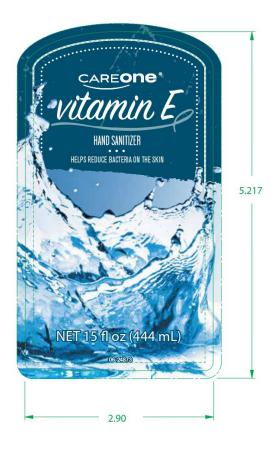




Front Label

**Back Label** 

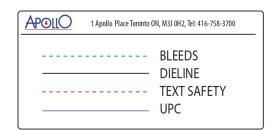






# **Front Label**

**Back Label** 

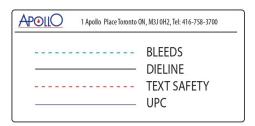






Front Label

**Back Label** 



# **CAREONE VITAMIN E HAND SANITIZER**

ethyl alcohol gel

Pro	duct	Information	
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:72476-200

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	650 mg in 1 mL		

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
POLYETHYLENE GLYCOL 300 (UNII: 5655G9 Y8 AQ)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
GLYCERIN (UNII: PDC6A3C0OX)				
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
.ALPHATO CO PHERO L ACETATE, DL- (UNII: WR1WPI7EW8)				
ACRYLATES/VINYL ISODECANO ATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8 MDB79 NA)				

P	Packaging				
#	# Item Code Package Description		Marketing Start Date	Marketing End Date	
1	NDC:72476-200- 02	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/06/2020		
2	NDC:72476-200- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/06/2020		
3	NDC:72476-200- 15	444 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/06/2020		
4	NDC:72476-200- 32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/06/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	10/06/2020		

# Labeler - Retail Business Services, LLC (967989935)

# **Registrant** - Apollo Health and Beauty Care (201901209)

Establishment					
Name	Address	ID/FEI	Business Operations		
Apollo Health and Beauty Care		20 19 0 120 9	manufacture(72476-200)		