# HAND SANITIZER- ethyl alcohol gel Walgreens

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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### Walgreens 469.000 469AA

#### **Active ingredient**

Ethyl alcohol 63%

#### **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. Incase 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 skin irritation or redness develops
- condition persists for more than 72 hours

### Keep out of reach of children

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

# **Directions**

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only 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, glycerin, tocopheryl acetate, retinyl palmitate, carbomer, fragrance, benzophenone-4, mannitol, cellulose, hydroxypropyl methylcellulose, ultamarines

### claims

Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds. Questions or comments? 1-800-925-4733

### adverse reactions section

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

walgreens.com

MADE IN U.S.A. WITH US AND FOREIGN COMPONENTS

# principal display panel

Well

Walgreens

Hand

Sanitizer

Coconut

water scent

Kills 99.99% of germs

2 FL OZ (59 mL)

### HAND SANITIZER

ethyl alcohol gel

Product	Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0469
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Route of Administration TOPICAL

# **Active Ingredient/Active Moiety**

Ingredient Name		Basis of Strength	Strength
ı	ALCOHOL (LINII) 3K9958V90M) (ALCOHOL - LINII) 3K9958V90M)	VI COHOI	5/15 mg in 1 ml

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)		
SULISOBENZONE (UNII: 1W6L629B4K)		
MANNITOL (UNII: 30WL53L36A)		
CELLULOSE ACETATE (UNII: 3J2P07GVB6)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
ULTRAMARINE BLUE (UNII: 139WR998BI)		

			Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
<b>1</b> N	NDC:0363-0469-16 5	9 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/26/2015			
Marketing Information						
ı	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final part333A		01/26/2015				

# Labeler - Walgreens (008965063)

# Registrant - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(0363-0469)

Revised: 6/2022 Walgreens