

## **HAND SANITIZER- ethyl alcohol gel**

**Walgreen**

*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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### **Hand Sanitizer**

**544**

#### **Active ingredient**

Ethyl alcohol 63%

#### **Purpose**

Antiseptic

#### **Use**

- 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 and redness develop
- 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, acrylates/C10-30 alkyl acrylate crosspolymer, benzophenone-4, mannitol, cellulose, hydroxypropyl methylcellulose, fragrance, yellow 5, blue 1, red 4, ultramarines

\*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

DISTRIBUTED BY: WALGREEN CO.  
200 WILMOT RD., DEERFIELD, IL60015  
100% satisfaction guaranteed  
walgreens.com  
544.000/544AA

## Principal display panel

Well at Walgreens

NEW

Hand Sanitizer Guava fruit

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:0363-0898

Route of Administration TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	567 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
glycerin (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
ULTRAMARINE BLUE (UNII: I39WR998BI)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0898-16	59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/04/2016	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/04/2016	

Labeler - Walgreen (008965063)

Registrant - Vi-Jon (790752542)

### Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(0363-0898)