# HAND SANITIZER- alcohol gel Hopkins Medical

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 ingredient**

Ethyl Alcohol 66.5%

#### **Purpose**

Antiseptic

#### Uses

- for handwashing to decrease bacteria on skin
- recommended for repeated use

#### Warnings

For external use only

## Flammable, keep away from fire or flame

**Do not use** in the eyes. If this happens, rinse thoroughly with water.

**Stop use and ask a doctor if** irritation and redness develop and persists for more than 72 hours

#### Keep out of reach of children

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

#### **Directions**

- wet hands thoroughly with product
- allow to dry without wiping
- children under 6 should be supervised while using this product

#### **Inactive ingredients**

aloe vera, carbomer, D&C green #5, D&C yellow #10, fragrance, purified water, triethanolamine

#### Principal Display Panel – 1.5 fl. oz. Bottle Label

Hopkins Waterless Hand Sanitizer

**Enriched with Aloe** 

**Hopkins** 

**Medical Products** 

Since 1945 1.5 FL OZ (46 ML) #697373

# **Hopkins** Waterless Hand Sanitizer

**Enriched with Aloe** 





1.5 FL OZ (46 ML) #697373

# HAND SANITIZER

alcohol gel

Product	Information
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:44662-1112

Route of Administration TOPICAL

# **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
alcohol (UNII: 3K9958V90M) (alcohol - UNII: 3K9958V90M)	alcohol	665 mL in 1 L

# **Inactive Ingredients**

Ingredient Name		
aloe vera leaf (UNII: ZY8 1Z8 3H0 X)		
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)		
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
water (UNII: 059QF0KO0R)		
trolamine (UNII: 9O3K93S3TK)		

### **Packaging**

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:44662-1112-0	0.046 L in 1 BOTTLE; Type 0: Not a Combination Product	08/26/2008	

# **Marketing Information**

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/26/2008	

# Labeler - Hopkins Medical (003080678)

# Registrant - Safetec of America, Inc. (874965262)

Establishment				
Name	Address	ID/FEI	Business Operations	
Safetec of America, Inc.		874965262	MANUFACTURE(44662-1112)	

Revised: 3/2020 Hopkins Medical