# ANTI BACTERIAL HAND SANITIZER- ethyl alcohol gel Dolgencorp Inc.

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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# Active Ingredients:

Ethyl Alcohol 62%

# **Purpose:**

Antiseptic

**Warnings:** For external use only.

Flammable. Keep away from fire or flame.

Do not apply around eyes.

Do not use in ears or mouth.

When using this product, avoid contact with eyes. In case of contact, flush eyes with water.

Stop use or ask for a doctor if redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children. Children must be supervised in use of this product.

#### Directions:

Pump as needed into your palms and thoroughly spread on both hands. Rub into skin until dry.

**Inactive ingredients:** Water, Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Aloe Barbadenis Leaf Juice

**Other information:** Do not store in temperature over 118 F.

**Uses:** Hand sanitizer to help decrease bacteria on the skin when water, soap & towel are not available. Recommended for repeated use.

## **Drug Facts**

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DOLGENCORP INC 100 MISSION,RIDGE GOODLETTSVILLE,TN 37072-2170 Anti Bacter

1 fl oz / 29 mL

Anti Bacterial Hand Sanitizer

MADE IN CHINA

8 10532 02728

# ANTI BACTERIAL HAND SANITIZER

ethyl alcohol gel

#### **Product Information**

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:55910-988

**Route of Administration** TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

#### **Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 903K93S3TK)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:55910-988- 29	29 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/31/2015		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	07/31/2015			

# Labeler - Dolgencorp Inc. (068331990)

Revised: 7/2015 Dolgencorp Inc.