

## **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%

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### **Purpose**

Antiseptic

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

Recommend for repeated use.

**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.

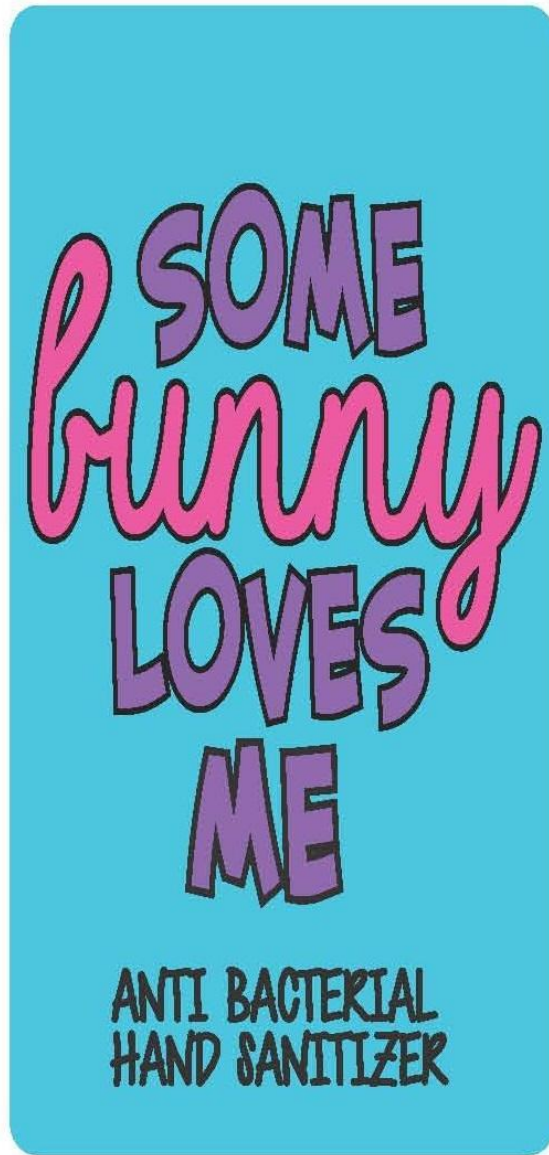
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**Directions:** Pump as needed into your palms and thoroughly spread on both hands. Rub into skin until dry.

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

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**INACTIVE INGREDIENTS:**Water, Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Aloe Barbadensis Leaf Juice



**Drug Facts**  
**Active ingredients..... Purpose**  
 Ethyl Alcohol 62%.....Antiseptic

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

1 fl oz / 29 mL

**MADE IN CHINA**



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<b>ANTI BACTERIAL HAND SANITIZER</b>			
ethyl alcohol gel			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:559 10-988
<b>Route of Administration</b>	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
<b>Ingredient Name</b>		<b>Basis of Strength</b>	<b>Strength</b>
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)		Alcohol	62 mL in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-988-08	29 mL in 1 BOTTLE, SPRAY		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/03/2014	

**Labeler** - Dolgencorp Inc (068331990)

Revised: 9/2014

Dolgencorp Inc