

**INSTANT HAND SANITIZER FRESH ALOE - ethyl alcohol gel  
BB17, LLC**

*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% v/v

**Purpose**

Antimicrobial

**KILLS MORE THAN 99.99% OF COMMON GERMS**

**WARNING:**

Flammable, Keep away from fire or flame. **FOR EXTERNAL USE ONLY. DO NOT USE IN THE EYES.**

DISCONTINUE USE IF IRRITATION AND REDNESS DEVELOP. IF CONDITION PERSISTS FOR MORE THAN 72 HOURS, CONSULT A DOCTOR.

**Keep out of reach of children.**

**DIRECTIONS:**

Rub into skin until dry.

**Other Information:**

Store below 110 F.

**INACTIVE INGREDIENTS:**Water, Aloe barbadenis, Leaf Juice, Glycerin,

Propylene Glycol, Carbomer, Triethanolamine, Fragrance, Tocopheryl Acetate, FD&C Blue NO.1



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**DRUG FACTS**

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**Inactive Ingredients:**  
Water, Aloe Barbadensis Leaf Juice, Glycerin, Propylene Glycol,

Carbomer,  
Triethanolamine,  
Fragrance, Tocopheryl Acetate, FD&C Yellow NO.5, FD&C Blue NO.1

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*FOR BEST USE, PLEASE SHAKE ME*

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E-mail: info@BB17usa.com

## INSTANT HAND SANITIZER FRESH ALOE

ethyl alcohol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:53603-1003
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	62 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMO POLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	

<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: HBR47K3TBD)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53603-1003-2	59.1 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	12/06/2012	

**Labeler** - BB17, LLC (828378294)

Revised: 12/2012

BB17, LLC