INSTANT HAND SANITIZER- 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.

Active Ingredient:

Ethyl Alcohol 62%

Purpose:

Antimicrobial

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 OR PHYSICIAN.

KEEP OUT OF REACH OF CHILDREN.

Directions:

Rub into hands until dry.

Water, Aloe barbadenis Leaf Juice, Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Fragrance, Tocopheryl Acetate, FD&C Red NO.33.

Other Information:

Store below 110 F.

KILLS MORE THAN 99.99% OF COMMON GERMS





Back Label - Peel off label

Note: Front side of back label.

WARNING: FLAMMABLE KEEP AWAY

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 OR PHYSICIAN.

FROM FIRE OR FLAME.

KEEP OUT OF REACH OF CHILDREN.

Note: This is the inside of a peel off label



KILLS MORE THAN 99.99% OF COMMON GERMS

DRUG FACTS
Active Ingredient:
Ethyl Alcohol 62%
Purpose:

Antimicrobial

Directions: Rub into hands until dry.

Other Information: Store below 110 ° F. **Inactive Ingredients:**

Water, Aloe
Barbadensis Leaf
Juice, Glycerin,
Propylene Glycol,
Carbomer,
Triethanolamine,
Fragrance,
Tocopheryl Acetate,
FD&C RED NO.33.

Distributed by BB17, LLC Cheyenne, WY 82001 All Rights Reserved Made in China E-mail: info@BB17usa.com

Note: Grey side sticks to bottle.

INSTANT HAND SANITIZER

ethyl alcohol gel

Droduct	Information
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Product Type HUMAN OTC DRUG In	Item Code (Source)	NDC:53603-1025
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Route of Administration 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

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Ingredient Name	Strength	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)		
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)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53603-1025-	29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/08/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/08/2015	

Labeler - BB17, LLC (828378294)

Revised: 6/2015 BB17, LLC