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:

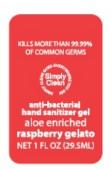
Rub into hands until dry.

Water, Aloe barbadenis Leaf Juice, Glycerin, Propylene Glycol, Olive Oil, Fragrance, Carbomer, Triethanolamine, D&C Red No.33.

Other Information:

Store below 110 F.

KILLS MORE THAN 99.99% OF COMMON GERMS





Note: This is the inside of a peel off label

WARNING:
FLAMMABLE KEEP AWAY
FROM FINE OR FLAME
FOR EXTERNAL USE ONLY.

Bull nite hands until dry.

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DO NOT USE IN THE EYES.
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JARTIATION AND REDNESS
DEVELOP. IF CONDITION
PERSISTS FOR MORE THAN
72 HOURS, CONSULT A
DOCTOR OR PHYSICIAN.

KEEP OUT OF REACH OF CHILDREN. Inactive Ingredients: Water, Aloe Barbadensis Leaf Juice, Glycerin, Propylene Glycol, Olive Oil, Fragrance, Carbomer, Triethanolamine, D&C Red No.33.

Other Information: Store below 110 ° F.

Hang Tag





INSTANT HAND SANITIZER

ethyl alcohol gel

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:53603-1048 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	
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)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
OLIVE OIL (UNII: 6UYK2W1W1E)	

l	Packaging				
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
l	1 NDC:53603-1048-1	29.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/16/20 17		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	0 1/16/20 17	

Labeler - BB17, LLC (828378294)

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