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:

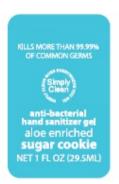
Rub into hands until dry.

Water, Aloe barbadenis Leaf Juice, Glycerin, Propylene Glycol, Olive Oil, Fragrance, Carbomer, Triethanolamine, FD&C Red No.4, FD&C Yellow No.5.

Other Information:

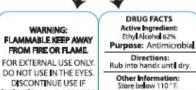
Store below 110 F.

KILLS MORE THAN 99.99% OF COMMON GERMS





Note: This is the inside of a peel off label



IRRITATION 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,

FD&C Red No.4, FD&C Yellow No.5.

Hang Tag





INSTANT HAND SANITIZER

ethyl alcohol gel

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53603-1049			
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: 9O3K93S3TK)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:53603-1049-1	29.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/16/20 17		
Marketing Information					
	Marketing Categor		Marketing Start Date	Marketing End Date	

Labeler - BB17, LLC (828378294)

Packaging

Revised: 1/2017 BB17, LLC