

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.

Inactive Ingredients:

Water, Aloe barbadenis Leaf Juice, Glycerin, Propylene Glycol, Fragrance, Carbomer, Triethanolamine, FD&C Yellow No.5, FD&C Blue No.1.

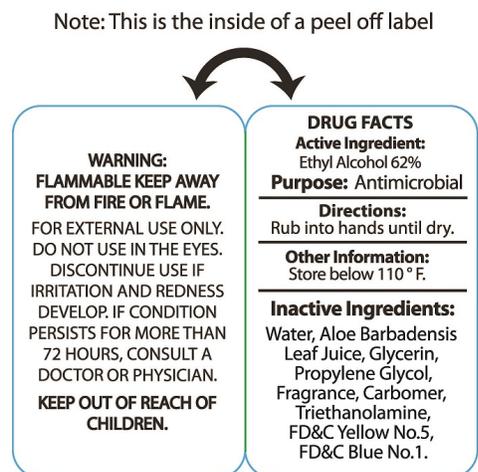
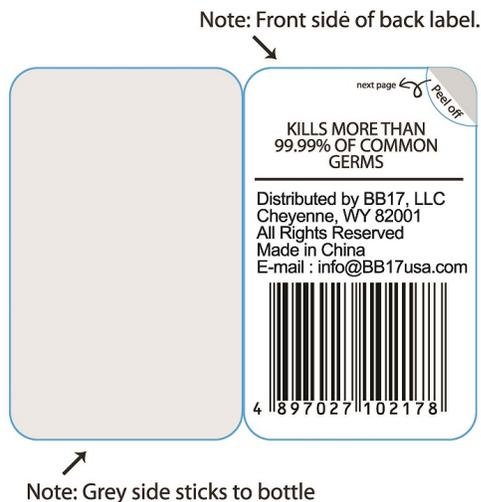
Other Information:

Store below 110 F.

KILLS MORE THAN 99.99% OF COMMON GERMS

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INSTANT HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:53603-1035

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: ZY81Z83H0X)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
	CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)			
	TROLAMINE (UNII: 9O3K93S3TK)			
	FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
	FD&C BLUE NO. 1 (UNII: HBR47K3TBD)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53603-1035-1	29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/26/2015	
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph not final	part333E	09/26/2015	

Labeler - BB17, LLC (828378294)