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, Fragrance, Carbomer, Triethanolamine, FD&C Blue No.1, FD&C Yellow No.5, FD&C Red No.4.

Other Information:

Store below 110 F.

KILLS MORE THAN 99.99% OF COMMON GERMS

Hand Tag Front



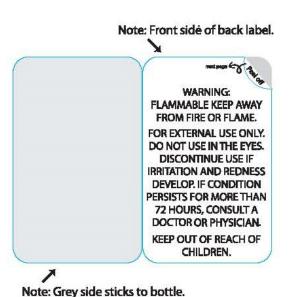
Hand Tag Back





5.5cm

Back Label - Peel off label



Inactive Ingredients: KILLS MORE THAN 99.99% OF COMMON GERMS Water, Aloe Barbadensis Leaf Juice, Glycerin, **DRUG FACTS** Propylene Glycol, Active Ingredient: Fragrance, Carborner, Ethyl Alcohol 62% Triethanolamine, FD&C Blue No.1, FD&C Yellow Purpose: No.5, FD&C Red No.4. **Antimicrobial** Directions: Rub into hands until dry. Distributed by BB17, LLC Cheyenne, WY 82001 All Rights Reserved Made in China E-mail: info@BB17usa.com Other Information: Store below 110 ° F.

Note: This is the inside of a peel off label

INSTANT HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53603-1027
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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
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 YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:53603-1027-	29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/21/2015	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	07/21/2015		

Labeler - BB17, LLC (828378294)

Revised: 1/2016 BB17, LLC