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

Anumicrobiai

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.

Other Information:

Store below 110 F.

KILLS MORE THAN 99.99% OF COMMON GERMS

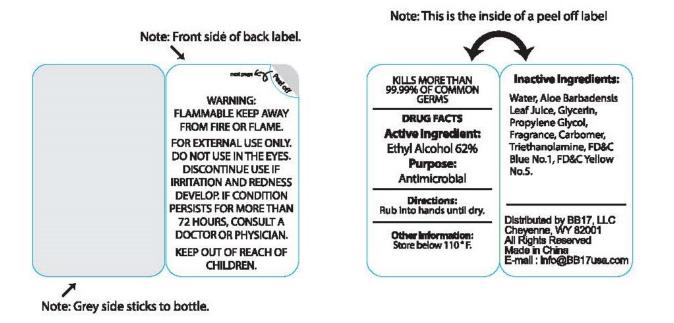
Hand Tag Front





5.5cm

Back Label - Peel off label



INSTANT HAND SANIT ethyl alcohol gel	IZER		
Product Information			
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:53603-1033
Route of Administration	TOPICAL		

icuve ingreuie	ent/Active Moiety		
	Ingredient Name	Basis of Streng	gth Strength
ALCOHOL (UNII: 3)	K9958V90M) (ALCOHOL - UNII:3K9958V90	M) ALCOHOL	62 mL in 100 mL
Inactive Ingred	lients		
	Strengtl		
FD&C BLUE NO. 1	(UNII: H3R47K3TBD)		
WATER (UNII: 059Q	PF0KO0R)		
ALOE VERA LEAF			
GLYCERIN (UNII: PI			
	C OL (UNII: 6DC9Q167V3)		
	POLYMER TYPE A (55000 CPS) (UNII: 59T	L3WG5CO)	
TROLAMINE (UNII:			
TROLAMINE (UNII.	9O3K93S3TK)		
	903K93S3TK) D. 5 (UNII: 1753WB2F1M)		
FD&C YELLOW NO	,		
FD&C YELLOW NO	,	Marketing Sta Date	rt Marketing En Date
FD&C YELLOW NO Packaging # Item Code	D. 5 (UNII: 1753WB2F1M)	Date	•
FD&C YELLOW NO Packaging # Item Code 1 NDC:53603-1033-	D. 5 (UNII: 1753WB2F1M) Package Description 29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a	Date	•
FD&C YELLOW NO Packaging # Item Code 1 NDC:53603-1033- 1	D. 5 (UNII: 1753WB2F1M) Package Description 29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Product	Date	•
FD&C YELLOW NO Packaging # Item Code 1 NDC:53603-1033-	D. 5 (UNII: 1753WB2F1M) Package Description 29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Product Product	Combination 07/21/2015	Date



Revised: 1/2016

15

BB17, LLC