HAND SANITIZER FRAGRANCE-FREE, PARABEN-FREE- ethyl alcohol gel Jocott Brands, Inc.

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.

HAND SANITIZER GEL WITH ALOE VERA - 65% Ethyl Alcohol

ACTIVE INGREDIENT

ETHYL ALCOHOL 65%

PURPOSE

ANTIMICROBIAL

USE

• HAND SANITIZER CAN HELP REDUCE BACTERIA ON THE SURFACE OF SKIN.

WARNINGS

FLAMMABLE. KEEP AWAY FROM FIRE OR FLAME.

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT DO NOT USE IN OR NEAR THE EYES. IN CASE OF CONTACT, RINSE EYES THOROUGHLY WITH WATER.

STOP USE AND ASK A DOCTOR IF IRRITATION OR RASH APPEARS AND LASTS.

KEEP OUT OF REACH OF CHILDREN. IF SWALLOWED GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

DIRECTIONS

- APPLY PRODUCT ALL OVER HANDS UNTIL DRY.
- CHILDREN SHOULD BE SUPERVISED AT ALL TIMES WHEN USING THIS PRODUCT.

OTHER INFORMATION

- STORE BELOW 110°F (43°C)
- MAY DISCOLOR CERTAIN FABRICS OR SURFACES

INACTIVE INGREDIENTS

PURIFIED WATER (AQUA), TOCOPHERYL ACETATE (VITAMIN E), ALOE BARBADENSIS (ALOE VERA) LEAF JUICE, CARBOMER, AMINOMETHYL PROPANOL, FRAGRANCE (PARFUM)







SIZE 3.436"W x 4.25"H
CORNERS: SQUARE
STOCK: WHITE BOPP
FINISH: MATTE
3"CORE / #4 POSITION

HAND SANITIZER FRAGRANCE-FREE, PARABEN-FREE

ethyl alcohol gel

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:78902-106 Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
GLYCERIN (UNII: PDC6A3C0OX)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)		
AMINO METHYLPRO PANOL (UNII: LU49 E6626Q)		

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:78902-106-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				
2	NDC:78902-106-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				
3	NDC:78902-106-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				
4	NDC:78902-106-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				
5	NDC:78902-106-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				
6	NDC:78902-106-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				
7	NDC:78902-106-24	739 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	06/12/2020			

Labeler - Jocott Brands, Inc. (080399826)

Revised: 7/2020 Jocott Brands, Inc.