PEARLESSENCE HAND SANITIZER HOLLYBERRY- 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.

PEARLESSENCE HAND SANITIZER GEL - WITH ALOE VERA - HOLLYBERRY - 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

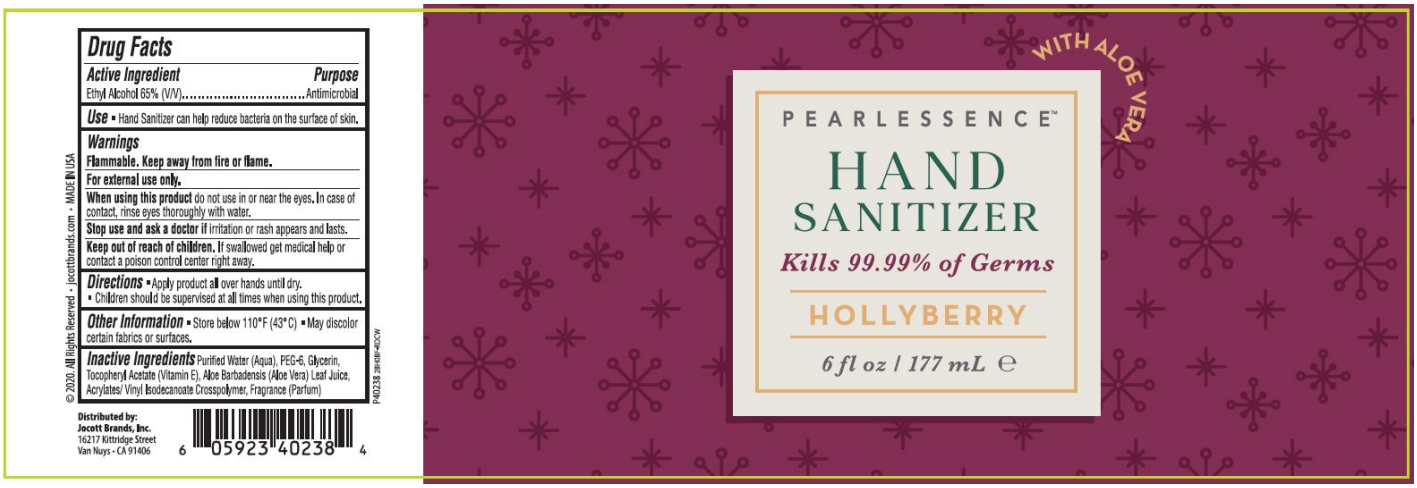
- APPLY PRODUCT ALL OVER HANDS UNTIL DRY.
- CHILDREN SHOULD BE SUPERVISED AT ALL TIMES WHEN USING THIS PRODUCT.

OTHER INFORMATION

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

INACTIVE INGREDIENTS

Purified Water (Aqua), PEG-6, Glycerin, Tocopheryl Acetate (Vitamin E), Aloe Barbadensis (Aloe Vera) Leaf Juice, Acrylates/Vinyl Isodecanoate Crosspolymer, Fragrance (Parfum)



PEARLESSENCE HAND SANITIZER HOLLYBERRY

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78902-117
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)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78902-117-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/01/2020	

Labeler - Jocott Brands, Inc. (080399826)

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