WARM VANILLA HAND SANITIZER- ethyl alcohol gel Greenbrier International, 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.

Warm Vanilla Hand Sanitizer

Active Ingredient Purpose

Ethyl Alcohol 62.0%.....Antimicrobial

Uses

- for handwashing to decrease the bacteria on the skin
- recommended for repeated use

□**Keep out of reach of children.**□ If swallowed, get medical help or contact a Poison Control Center right away.

Other information

- store at 20oC to 25oC (68o to 77oF)
- may discolor certain fabrics

□ Warnings

For external use only.

Flammable, keep away from heat and flame.

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Directions

- wet hands thoroughly with product
- briskly rub hands together until dry
- supervise children under 6 years in the use of this product

Inactive Ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, fd&c yellow no.5, fd&c red no.33.



DISTRIBUTED BY: GREENBRIER INTERNATIONAL, INC. **500 VOLVO PARKWAY**, CHESAPEAKE, VA 23320 **MADE IN CHINA**

Drug Facts

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143588-14454-002-1306



WARM VANILLA HAND SANITIZER

ethyl alcohol gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:33992-3004		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	147 mL in 237 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)				
TROLAMINE (UNII: 903K93S3TK)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				

l	Packaging				
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 N	DC:33992-3004-1	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/24/2013	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	05/24/2013		

Labeler - Greenbrier International, Inc. (610322518)

Establishment				
Name	Address	ID/FEI	Business Operations	
China Ningbo Shangge Cosmetic Technology Corp.		529287434	manufacture(33992-3004)	

Revised: 4/2019 Greenbrier International, Inc.