FOLLOW THE RAINBOW ANTIBACTERIAL HAND SANITIZER SWEET BERRIES SCENTED- alcohol gel

Ganzhou Olivee Cosmetic Co., Ltd.

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

Drug Facts

Active ingredient

Alcohol 65%

Purpose

Antiseptic

USE

Decrease bacteria on hands

Warnings

For external use only.

When using this product

avoid contact with eyes; in case of contact, flush eyes with water.

Stop use and ask a doctor if

• irritation and redness develops and persists.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Wet hands thoroughly with product and allow to dry.

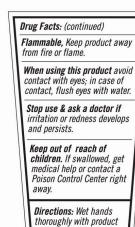
Inactive ingredients

Water (Aqua), Acrylates/c 10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrace (Parfum). May contain: FD&C No. 4 (C.l. 14700), D&C Red No. 33 (Cl 17200), FD&C Yellow 5, FD&C Blue No. 1 (Cl 42090).

package







and allow to dry.



Distributed by Best Accessory Group 65 West 36th St. 6th Fl. New York, NY 10018 Rue de la Victoire 26 Suite: 132494, 1060 Brussels, Belgium Designed in USA · Made in China Item#: TJ1602

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alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:56136-093

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (ALCOHOL - UNII:3K9958V90M)

Inactive Ingredients			
Ingredient Name	Strength		
water (UNII: 059QF0KO0R)			
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)			
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			
GLYCERIN (UNII: PDC6A3C0OX)			
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
D&C RED NO. 33 (UNII: 9DBA0SBB0L)			

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:56136-093-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2020				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	07/30/2020				

Labeler - Ganzhou Olivee Cosmetic Co., Ltd. (543008195)

Establishment					
Name	Address	ID/FEI	Business Operations		
Ganzhou Olivee Cosmetic Co., Ltd.		543008195	manufacture(56136-093)		

Revised: 7/2020 Ganzhou Olivee Cosmetic Co., Ltd.