WHIPPED VANILLA HAND SANITIZER- 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 66%

Purpose

Antiseptic

$\Box Use$

Decreases bacteria on hands

Warnings

For external use only.

Flammable. Keep product away from heat or flame.

Do not use: I in children less than 2 months of age. On open skin wounds

When using this product Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash develops.

Questions:

1-866-983-8582

Keep out of reach of children.

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

Directions

Apply sanitizer to hands. Rub hands together until dry.

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance, Water.

Packaging



Cleanse and condition hands without the use of water.

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Warnings For external use only Flammable: Keep away from heat or flame. Do not use . in children less than 2 months of age • on open skin wounds • When using

DRUG FACTS CONTINUED ▶

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DRUG FACTS CONTINUED >

Drug Facts (continued)

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NOT TESTED ON ANIMALS MADE IN CHINA DISTRIBUTED BY ULTA INC. BOLINGBROOK, IL 60440 ULTA.COM

#UltaBeautyCollection

WHIPPED VANILLA HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:56136-342

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 66 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)			
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56136-342-01	29 mL in 1 BOTTLE: Type 0: Not a Combination Product	06/27/2020	

Marketing Information

0			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/27/2020	

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

Establishment							
Name	Address	ID/FEI	Business Operations				
Ganzhou Olivee Cosmetic Co., Ltd.		543008195	manufacture (56 136 - 342)				

Revised: 6/2020 Ganzhou Olivee Cosmetic Co., Ltd.