YOU ARE A MAGICAL UNICORN ANTIBACTERIAL HAND SANITIZER WATERMELON 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

Ethyl Alcohol 65%

Purpose

Antiseptic

Use

Decrease bacteria on hands.

Warnings

For external use only.

Flammable. Keep product away from fire or flame.

When using this product

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

Stop use and ask a doctor if

irritation or 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/C10-30 ALKYL ACRYLATE CROSSCOPOLYMER, AMINOMETHYL PROPANOL, Fragrance (Parfum), May contain: D&C Red No. 33 (Cl 17200), FD&C Red No. 4 (Cl 14700), FD&C Yellow No. 5 (Cl 19140), FD&C Blue No. 1 (Cl 42090).

No Animal Testing

Packaging

YOU ARE A MAGICAL UNICORN ANTIBACTERIAL HAND SANITIZER WATERMELON SCENTED

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:56136-429
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)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Other Ingredients		
Ingredient Kind	**Ingredient Name**	**Quantity**
May contain	**D&C RED NO. 33** (UNII: 9DBA0SBB0L)	
May contain	**FD&C RED NO. 4** (UNII: X3W0AM1JLX)	
May contain	**FD&C YELLOW NO. 5** (UNII: I753WB2F1M)	
May contain	**FD&C BLUE NO. 1** (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56136-429-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/18/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
Ganzhou Olivee Cosmetic Co., Ltd.		543008195	manufacture(56136-429)

Revised: 10/2020 Ganzhou Olivee Cosmetic Co., Ltd.