

**YOU ARE MY SUNSHINE ANTIBACTERIAL HAND WASH WATERMELON SCENTED-
benzalkonium chloride liquid
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 Wash

Drug Facts

Active ingredient

Benzalkonium Chloride 0.1500%

Purpose

Antibacterial

USE

Helps eliminate 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 or redness develops and persists.

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

Directions

Apply onto wet hands working into a lather. Rinse thoroughly.

Inactive ingredients

Water (Aqua), Cocamidopropyl Betaine, Decyl Glucoside, Cocamidopropylamine Oxide Peg-80 Glyceryl Cocoate, Fragrance (Parfum), Phenoxyethanol, Benzyl Alcohol, Peg-15 Distearate, Glycol Distearate, Sodium Chloride, Citric Acid. May contain: FD &C Red No. 4 (C.I. 14700), FD&C Yellow No. 5 (C.I. 19140), FD&C Blue No. 1 (C.I. 42090), FD&C Red No.33 (C.I. 17200).

Hand wash



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benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56 136-513
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
GLYCERIN (UNII: PDC6A3C0OX)	
ACETYLTRIETHYL CITRATE (UNII: 5WBR36T90E)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PEG-80 GLYCERYL COCOATE (UNII: A7D04GUR0Z)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
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)	
May contain	D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56136-513-01	350 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/27/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
Ganzhou Olivee Cosmetic Co., Ltd.		543008195	manufacture(56136-513)

Revised: 11/2020

Ganzhou Olivee Cosmetic Co., Ltd.