

HAND SANITIZER FRESH KIWI 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

Decreases 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.

Choking Hazard

Contains small parts. Not for children under 3 years.

Keep out of reach of children.

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

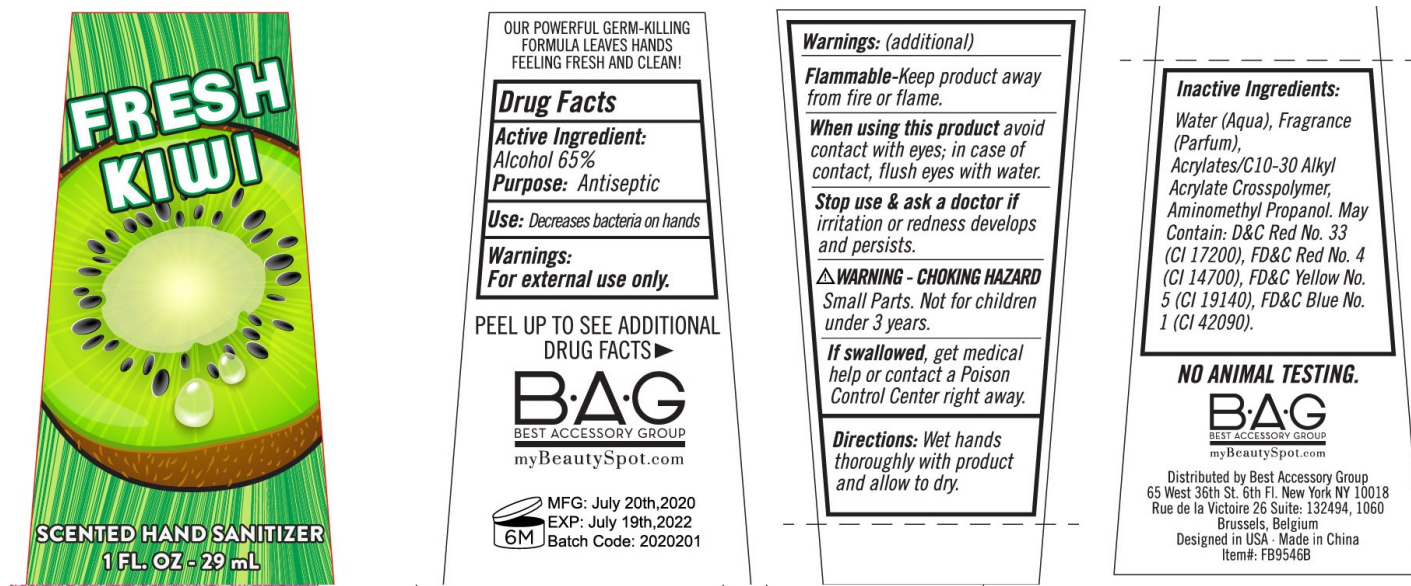
Directions

Wet hands thoroughly with product and allow to dry.

Inactive ingredients

Water (Aqua), Fragrance (Parfum), Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol. May contain: D&C Red No. 33 (CI 17200), FD&C Red No. 4 (CI 14700), FD&C Yellow No. 5 (CI 19140), FD&C Blue No. 1 (CI 42090).

Packaging



HAND SANITIZER FRESH KIWI SCENTED

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56 136 -339
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)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WP1EW8)	

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
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1	NDC:56136-339-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333E		06/25/2020	

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

Establishment			
Name	Address	ID/FEI	Business Operations
Ganzhou Olivee Cosmetic Co., Ltd.		543008195	manufacture(56136-339)