WAVE HAND SANITIZER- alcohol gel C & A MARKETING, INC.

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.

CA Global AO Wave Sanitizer Gel 75

Active Ingredient

Ethyl Alcohol 75%v/v

Purpose

Antiseptic

Uses

For reducing bacteria on the skin.

Warnings

For external use only. Flammable. Keep away from heat or flame

When using this product

Avoid contact with eyes, and if eye contact occurs, flush thoroughly with water and seek medical advice

Stop use and ask a doctor

if irritation or rash occurs. These may be the signs of a serious condition.

Keep out of reach of children

In case of accidentail ingestion, contact a doctor or Poison Control Center immediately.

Directions

• Dispense into dry hands and rub until dry. • Children under 6 years of age should be supervised by an adult when using this product.

Other information

• Store below 110°F (43°C) • May discolor certain fabrics or surfaces.

Inactive ingredients

Carbomer, water

Package Label

ANTIBACTERIAL HAND SANITIZER

Drug Facts

Active ingredient Purpose Ethyl Alcohol 75% Antiseptic

Uses

For reducing bacteria on skin

Warnings For external use only

Flammable

Keep away from fire and heat

When using this product

Avoid contact with eyes, and if eye contact occurs, flush thoroughly with water and seek medical advice

Stop use and ask a doctor if

Keep out of reach of children

Directions

· Dispense into dry hands and rub until dry Children under 6 years of age should be supervised by an adult when using this product

Other information

•Store below 110°F (43°C)
•May discolor certain fabrics or surfaces

Inactive ingredients

Carbomer, wate





75% ALCOHOL



16.9 oz./500 ml



- KILLS 99.9% OF GERMS
- FRAGRANCE-FREE
- DRIES OUICKLY

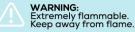
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C&A Marketing, Inc. wavessentials.com For sales inquiries:

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WAVE HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78754-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 75 mL in 100 mL

Inactive Ingredients

Strength Ingredient Name WATER (UNII: 059QF0KO0R) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)

Packaging

П	i uckug mg					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:78754-002- 08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020			
	2 NDC:78754-002- 16	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020			

3	NDC:78754-002- 28	3790 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020	08/28/2020
4	NDC:78754-002- 68	5000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/28/2020	
5	NDC:78754-002- 02	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/06/2020	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	06/15/2020				

Labeler - C & A MARKET ING, INC. (134723316)

Revised: 9/2020 C & A MARKETING, INC.