

**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 Wave Sanitizer Gel 75

Active Ingredient

Alcohol 75%v/v

Purpose

Antiseptic

Uses

- Hand Sanitizer to help reduce bacteria on the skin. For use when soap and water are not available

Warnings

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

Do not use

- on children less than 2 months of age ■ on open skin wounds

When using this product

keep out of eyes, ears, ears and mouth. In case of eye contact immediately flush eyes thoroughly with water

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

- Place enough product on hands to cover all surfaces. Rub hands together until dry. ■ Supervise children under 6 years of age when using this product to avoid swallowing.

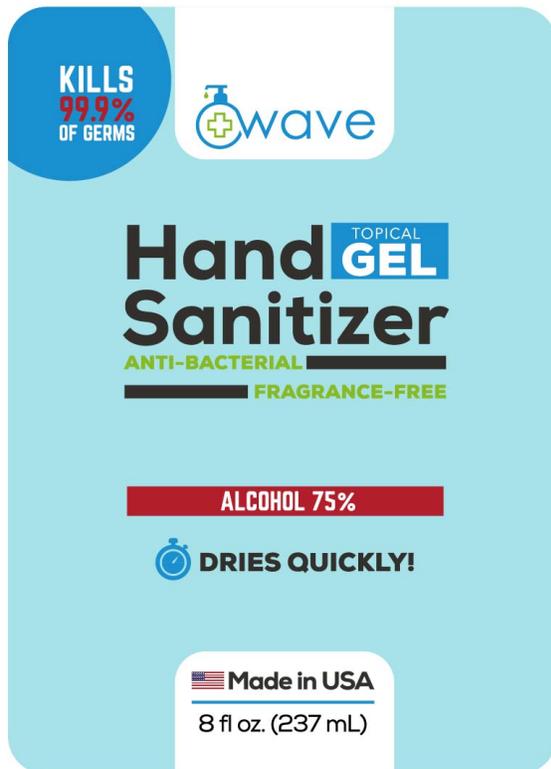
Other information

- Store between 15°-30°C (59°-89°F) ■ Avoid freezing and excessive heat above 40°C (104°F) ■ May discolor fabrics or surfaces.

Inactive ingredients

- Glycerin, hydroxypropyl cellulose, water

Package Label



WAVE HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78754-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78754-001-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020	
2	NDC:78754-001-16	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020	
3	NDC:78754-001-28	3790 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020	

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 MARKETING, INC. (134723316)

Revised: 6/2020

C & A MARKETING, INC.