

HAND SANITIZER GEL- alcohol gel

Resource One 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.

Hand Sanitizer Gel

Drug Facts

Active Ingredient:

Ethyl Alcohol 70%

Purpose:

Antiseptic

Uses:

Hand sanitizer to help reduce the bacteria that potentially will cause disease

Warnings:

For External Use Only

Flammable keep away from children

When using this product:

Avoid contact with face, eyes, and broken skin. If eye contact occurs, flush thoroughly with water and seek medical advise.

Stop and ask a doctor if:

Irritation and redness develops

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 rub into skin until dry.

Children under 6 years of age should be supervised by an adult when using this product

Inactive Ingredients:

Glycerin, Carbomer

Package Labeling: 946 ml

Description

Hand sanitizer is a Gel 70% alcohol hand sanitizer designed to be used in a dispenser or table top pump bottle.

Directions for Use

Hands should be cleansed of visible soils with soap and water. Apply small amount of product, rubbing hands thoroughly with product until dry. No rinsing needed.



Gator Chemical
2202 Industrial Blvd
Sarasota, FL 34234
www.gatorchemical.com
800-224-9199

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Emergency Telephone (24Hr)
1-352-323-2500 (International)
1800-535-5053 (North America)

Package Labeling:3785ml

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HAND SANITIZER GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77188-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77188-002-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/29/2020	
2	NDC:77188-002-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/29/2020	

Labeler - Resource One Inc (081366510)

Revised: 11/2020

Resource One Inc