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

Package Labeling:3785ml

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 HOMOPOLYMER, 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)

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