HAND SANITIZER- alcohol gel GoodFibers

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.

Alcohol

Active Ingredient

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

Uses

- Hand sanitizer to decrease bacteria on the skin
- Recommended for repeated use

Warnings

Warnings

For external use only Flammable

• Keep away from fire or flame

When using

When using this product:

• Do not use in or near eyes

If contact occurs, rinse eyes thoroughly with water.

Stop use

Stop use and ask a doctor

• If irritation and redness persist for more than 72 hours.

Keep out of reach

Keep out of reach of children

In case of accidental ingestion, seek medical attention or contact a poison control centerimmediately

Directions

Directions

- Place enough product on your palm to thoroughly spread on both hands and rub into the skin until dry.
- Children under six years of age should be supervised when using this product.

Other information

Other Information

• Store at 1 °C - 30 °C (34 °F - 86 °F)

Inactive ingredients

Inactive Ingredients

Carbomer, Glycerin, Triethanolamine, Water

PDP







Ethyl Alcohol

70%

all clear

Hand

Sanitizer

Fragrance Free

Kills 99.99 of germs

16.9 fl oz 500ml

HAND SANITIZER

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77070-700	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
ACRYLIC ACID (UNII: J94PBK7X8S)	

Product Characteristics				
Color	white (Clear colorless)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

ı	Packaging				
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:77070-700-80	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/07/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	05/07/2020		

Labeler - GoodFibers (105870470)

Registrant - GoodFibers (105870470)

Establishment				
Name	Address	ID/FEI	Business Operations	
Biotechnologia Educacion y Genetica, S.A. de C.V.		951580322	manufacture(77070-700)	

Revised: 7/2020 GoodFibers