

HAND SANITIZER- alcohol gel
Doctor Fashion SAPI de CV

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

Drug Facts

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

To decrease bacteria on the skin when water, soap & towel are not available – Recommended for repeated use.

Warnings

For external use only.

Flammable — Keep away from fire or flame.

When using this product keep out of eyes. In case of contact with eyes, rinse thoroughly with water. Do not use on broken or irritated skin.

Stop use and ask a doctor if irritation or redness develop and last more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Apply enough product to wet hands. Rub hands together until dry. Supervise children in use of this product.

Other information

- Do not store above 105 °F (40 °C).
- May discolor certain wood surfaces.

Inactive Ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, *aloe barbadensis* leaf juice.

PRINCIPAL DISPLAY PANEL - 200 ml Bottle Label

HAND
SANITIZER

CONTAINS

70%

ETHYL

ALCOHOL

TOPICAL

SOLUTION

WITH

ALOE VERA

6.7 Fl. Oz. (200 ml)

HAND SANITIZER

CONTAINS

**70% ETHYL
ALCOHOL**



**TOPICAL
SOLUTION**



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Made in México. Distributed by:



HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75625-001
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: 059QF0K00R)		
		CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
		trolamine (UNII: 9O3K93S3TK)		
		glycerin (UNII: PDC6A3C0OX)		
		propylene glycol (UNII: 6DC9Q167V3)		
		aloe vera leaf (UNII: ZY81Z83H0X)		
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75625-001-01	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2020	
2	NDC:75625-001-02	415 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL		part333A	02/05/2020	

Labeler - Doctor Fashion SAPI de CV (951576481)

Establishment

Name	Address	ID/FEI	Business Operations
Absara Cosmetics SAPI de CV		816161236	MANUFACTURE(75625-001)

Establishment

Name	Address	ID/FEI	Business Operations
TECNOGLOBAL PH7		813006665	MANUFACTURE(75625-001)

Revised: 4/2020

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