

HAND SANITIZER- alcohol gel
GIA EVOLUTION SA 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.

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

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

Do not use

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

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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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 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

purified water USP, hypromellose.

Package Label - Principal Display Panel

20000000 mL NDC: 77231-050-09

Alcohol Antiseptic 70%
gel

Hand Sanitizer
Non-sterile gel

(20,000 L)

Manufactured by:
Gia Evolution S.A. de C.V.
Carr. Aguascalientes-Zacatecas Km. 8.
Aguascalientes, AGS. México. 20110
+52 (449) 158 3500

Drug Facts	
Active ingredient[s]	Purpose
Alcohol 70% v/v.....	Antiseptic
Use[s]	
Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.	
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<ul style="list-style-type: none"> ▪ 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	
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Inactive ingredients	
Hypromellose, purified water USP.	

3875 mL NDC: 77231-050-06

Alcohol Antiseptic 70%
gel

Hand Sanitizer
Non-sterile gel

1 gal (3.785 L)

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Drug Facts	
Active ingredient[s]	Purpose
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Other information	
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Inactive ingredients	
Hypromellose, purified water USP.	

946.35 mL NDC: 77231-050-05

Alcohol Antiseptic 70%
gel

Hand Sanitizer
Non-sterile gel

1 Gal oz (3.875 L)

Manufactured by:
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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 1-30C (59-85F) • Avoid freezing and excessive heat above 40C (104F)	
Inactive ingredients Hydroxyethylcellulose, purified water USP.	

500 mL NDC: 77231-050-04

Alcohol Antiseptic 70%
gel

Hand Sanitizer
Non-sterile gel

16.9 oz (500 mL)

Manufactured by:
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Drug Facts	
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Inactive ingredients Hydroxyethylcellulose, purified water USP.	

236.59 mL NDC: 77231-050-03

Alcohol Antiseptic 70%
gel

Hand Sanitizer
Non-sterile gel

8 oz (236.6 mL)

Manufactured by:
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Other information • Store between 1-30C (59-85F) • Avoid freezing and excessive heat above 40C (104F)	
Inactive ingredients Hydroxyethylcellulose, purified water USP.	

189.27 mL NDC: 77231-050-10

Alcohol Antiseptic 70%
gel

Hand Sanitizer
Non-sterile gel

6.4 oz (189.3 mL)

Manufactured by:
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Drug Facts	
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Inactive ingredients Hydroxyethylcellulose, purified water USP.	

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77231-050
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
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	0.7 g in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77231-050-09	20000000 mL in 1 TANK; Type 0: Not a Combination Product	03/30/2020	
2	NDC:77231-050-06	3875 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
3	NDC:77231-050-05	946.35 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:77231-050-04	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:77231-050-03	236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:77231-050-10	189.27 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - GIA EVOLUTION SA DE CV (951577293)

Registrant - GIA EVOLUTION SA DE CV (951577293)

Establishment

Name	Address	ID/FEI	Business Operations
GIA EVOLUTION SA DE CV		951577293	manufacture(77231-050) , pack(77231-050) , label(77231-050)

Revised: 6/2020

GIA EVOLUTION SA DE CV