

HAND SANITIZER- alcohol gel

Diquisa, S.A. De C.V.

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

Active Ingredient(s)

Ethyl Alcohol 70% v/v.

Purpose

Antiseptic

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

- on 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.

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

Inactive ingredients

Carbomer, Glycerin, Purified water, Trolamine

Package Label - Principal Display Panel

3780 mL NDC: 78361-003-01

AVOS
ANTIBACTERIAL
**HAND
SANITIZER**

ALCOHOL 70%
FRAGRANCE FREE
GENTLE ON SKIN · QUICK DRY

128 FL OZ (1 GAL)

1000 mL NDC: 78361-003-02

Drug Facts

Active ingredients	Purpose
Ethyl alcohol 70%.....	Antiseptic

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Inactive ingredients

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Manufactured by:
DIQUIISA, S.A. DE C.V.
Monterrey, NL 64630
www.avosfamily.com
Contact:
+52 (81) 8333-0570
MADE IN MEXICO



AVOS

ANTIBACTERIAL

HAND SANITIZER



*ALCOHOL 70%
FRAGRANCE FREE
GENTLE ON SKIN · QUICK DRY*

33.8 FL OZ (1 L)

Drug Facts

Active ingredients

Ethyl alcohol 70%.....Antiseptic

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500 mL NDC: 78361-003-03

AVOS

ANTIBACTERIAL

HAND SANITIZER



*ALCOHOL 70%
FRAGRANCE FREE
GENTLE ON SKIN · QUICK DRY*

16.9 FL OZ (500 ML)

Drug Facts

Active ingredients

Ethyl alcohol 70%.....Antiseptic

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250 mL NDC: 78361-003-04

AVOS

ANTIBACTERIAL

HAND SANITIZER



*ALCOHOL 70%
FRAGRANCE FREE
GENTLE ON SKIN · QUICK DRY*

8.45 FL OZ (250 ML)

Drug Facts

Active ingredients

Ethyl alcohol 70%.....Antiseptic

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330 mL NDC: 78361-003-05

AVOS

ANTIBACTERIAL

**HAND
SANITIZER**

ALCOHOL 70%

FRAGRANCE FREE

GENTLE ON SKIN · QUICK DRY

11.16 FL OZ (330 ML)



Drug Facts

Active ingredients

Ethyl alcohol 70%.....Antiseptic

Purpose

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

Warnings

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Inactive ingredients

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NDC 78361-003-05



60 mL NDC: 78361-003-06

AVOS

ANTIBACTERIAL

HAND SANITIZER

ALCOHOL 70%
FRAGRANCE FREE
GENTLE ON SKIN · QUICK DRY

2 FL OZ (60 ML)

Drug Facts

Active ingredients

Ethyl alcohol 70%.....Antiseptic

Purpose

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

Warnings

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Inactive ingredients

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NDC 78361-003-06

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78361-003
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
TROLAMINE (UNII: 9O3K93S3TK)	0.1 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.4 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	0.5 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78361-003-01	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
2	NDC:78361-003-02	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
3	NDC:78361-003-03	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
4	NDC:78361-003-04	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
5	NDC:78361-003-05	330 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
6	NDC:78361-003-06	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/2020	

Labeler - Diquisa, S.A. De C.V. (815736509)

Establishment

Name	Address	ID/FEI	Business Operations
Diquisa, S.A. De C.V.		815736509	manufacture(78361-003)

Revised: 8/2020

Diquisa, S.A. De C.V.