

**HAND SANITIZER- isopropyl alcohol gel**  
**DEWARS COSMETIQUE LTDA**

*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**

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Isopropyl Alcohol (75%, v/v) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

**Active Ingredient(s)**

Isopropyl 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.

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

### Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

### Package Label - Principal Display Panel

Drug Facts	
Active Ingredient.....	Purpose
Ethyl Alcohol 70%.....	Antiseptic
<b>Inactive Ingredients:</b>	
• Water(Aqua), Glycerin, Carbomer, Triethanolamine, Tocopheryl acetate, Aloe Barbadosis, Leaf extract.	
<b>Warnings: Flammable. Keep away from fire or flame.</b>	
For external use only. When using this product avoid contact with eyes. <b>Keep out of reach of children.</b> If is swallowed get medical help promptly.	
<b>Stop use and ask a doctor:</b> If any irritation appears, these may be signs of a serious condition.	
<b>Uses:</b>	
• To help decrease bacteria on the skin. • Recommended for repeated use.	
<b>Directions:</b>	
• Place a small amount of the product in one hand and rub hands together even between fingers until dry. • Supervise children under 6 year of age when using this product to avoid swallowing.	
<b>Other Information:</b>	
Store in a cool place below 110°F (43° C). Avoid freezing. May discolor certain fabrics of surfaces.	

16.91 oz (500 ml)

250 ml NDC: 79180-001-50

<b>HAND SANITIZER</b>
isopropyl alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79180-001
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>CHLORHEXIDINE GLUCONATE</b> (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	0.25 g in 100 mL
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>ALOE ARBORESCENS LEAF</b> (UNII: 09TD8L5SQV)	
<b>CASTOR OIL</b> (UNII: D5340Y2I9G)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>2-PHENYLPROPANAL PROPYLENE GLYCOL ACETAL</b> (UNII: 1ZRR9A405A)	
<b>CARBOBENZOXYCARBONYL-PHENYLALANYL-ALANINYLDIAZOMETHANE</b> (UNII: FQ11Y48OCY)	
<b>.ALPHA.-TOCOPHEROL ACETATE, D-</b> (UNII: A7E6112E4N)	

**Product Characteristics**

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:79180-001-45	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
2	NDC:79180-001-50	16.91 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	07/08/2020
3	NDC:79180-001-84	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
4	NDC:79180-001-55	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2020	



### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/25/2020	

**Labeler** - DEWARS COSMETIQUE LTDA (882315797)

Revised: 7/2020

DEWARS COSMETIQUE LTDA