## HAND SANITIZER- alcohol gel COMERCIALIZADORA Y MULTISERVICIOS CAMALDO 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.

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#### **RESPOND 1.69 OZ LABEL**

## **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**

## Package Label - Principal Display Panel



Drug Facts
Active Ingradients
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Ethyl Abohol 70%.

Antiaeptic
Utes:
Hand somitizer to help reduce bacteria on skin that could dause disease, keeps hands clean for up to 8 hours
Warnings: Filammable keep away from the or florme For external use only
When using this product do not use in or near eyes, in case of contact rinse eyes thoroughly with water.
Shap and eak elector if initiation or rosh appears and joints
Keep out of reach of children if swellowed seek medical help or contact a Palson Control Center right away
Directions:
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1.69 OZ (50ml)

50 mL NDC:79479-001-50

## HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79479-002
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	
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)		
WATER (UNII: 059QF0KO0R)		
CARBO MER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM30 7FC)		
ZINC (UNII: J41CSQ7QDS)		
TROLAMINE (UNII: 903K93S3TK)		
ALOE VERA LEAF (UNII: ZY81Z83H0 X)		

I	Packaging				
I	#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
I	1 N	DC:79479-002-50	50 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

5 5	* *	 	9
OTC monograph not final	part333A	07/08/2020	

# Labeler - COMERCIALIZADORA Y MULTISERVICIOS CAMALDO SA DE CV (951580121)

Revised: 7/2020 COMERCIALIZADORA Y MULTISERVICIOS CAMALDO SA DE CV