# AMSINO HAND SANITIZER- ethyl alcohol gel MCCORD RESEARCH

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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#### AMSINO HAND SANITIZER

# Active Ingredient

Ethyl Alcohol

### **Purpose**

**Antiseptic** 

#### Uses

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 produc**t 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 15-30°C (59-86F)

Avoid freezing and excessive heat above 40°C (104F)

# Inactive Ingredients

Water, Fragrance, Carbomer, PEG-40 Hydrogenated Castor Oil, chlorphenesin, Aminomethyl Propanol

#### **Product Label**



# **AMSINO HAND SANITIZER**

ethyl alcohol gel

Product Information

**Route of Administration** 

1 Todace Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71358-020

TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	620 mg in 1 mL
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Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)		
CHLORPHENESIN (UNII: 1670DAL4SZ)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:71358-020-	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/07/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/07/2020	

# Labeler - MCCORD RESEARCH (010011284)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Wasatch Product Development, LLC		962452533	manufacture(71358-020)	

Revised: 1/2022 MCCORD RESEARCH