

**HAND SANITIZER- ethyl alcohol gel**  
**MINISO DEPOT CA, INC.**

*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  
Ethyl Alcohol 62%

Purpose  
Antiseptic

**Uses** Decrease bacteria on hands

**Warnings** For external use only.

Keep out of eyes. In case of contact with eyes, flush

Keep it out of reach of children.

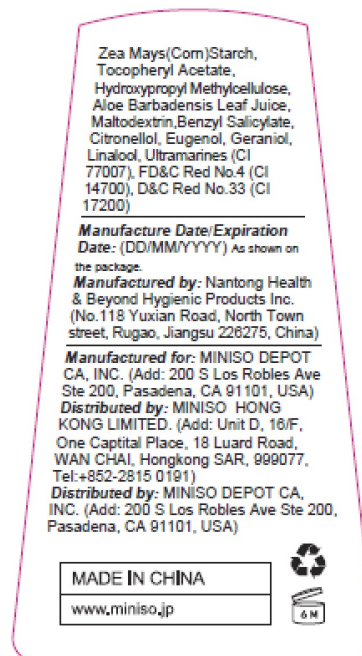
**Stop use and ask a doctor if** irritation or redness develop.If swallowed,get medical help or contact a doctor right away.

**Storage:** Flammable,Avoid direct sunlight and keep away from fire.

**Directions:** Wet hands thoroughly with product and allow to dry without wiping.

**Inactive Ingredients:**

Water (Aqua),Glycerin, Propylene Glycol,Fragrance (Parfum), Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Lactose, Microcrystalline Cellulose, Sucrose, Zea Mays(Corn)Starch, Tocopheryl Acetate, Hydroxypropyl Methylcellulose, Aloe Barbadensis Leaf Juice, Maltodextrin,Benzyl Salicylate, Citronellol, Eugenol, Geraniol, Linalool, Ultramarines (CI77007), FD&C Red No.4 (CI14700), D&C Red No.33 (CI17200)



**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73950-001
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
GERANIOL (UNII: L837108USY)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
ULTRAMARINE BLUE (UNII: I39WR998B1)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
.ALPHA.-LACTOSE (UNII: MJF4JAT10B)	
SUCROSE (UNII: C151H8M554)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
STARCH, CORN (UNII: O8232NY3SJ)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
EUGENOL (UNII: 3T8H1794QW)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73950-001-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/23/2020	

