

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 thoroughly with water.

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, Aloe Barbadensis Leaf Juice, Glycerin, Propylene Glycol, Fragrance, Acrylates/C10-30 AlkyI Acrylate Crosspolymer, Aminomethyl Propanol, Maltodextrin.



HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73950-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73950-005-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/26/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/26/2020	

Labeler - MINISO DEPOT CA, INC. (056958961)

Revised: 5/2020

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