MIXLINE- alcohol solution
EMPERO LLC

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

Alcohol 75% v/v

Purpose

Antiseptic

Uses

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

Flammable. Keep away from heat or flame. For external use only.

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.

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

Alcohol, Water, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine, Aloe Barbadensis Leaf Juice.

Principal Display Panel

NDC 76778-075-17

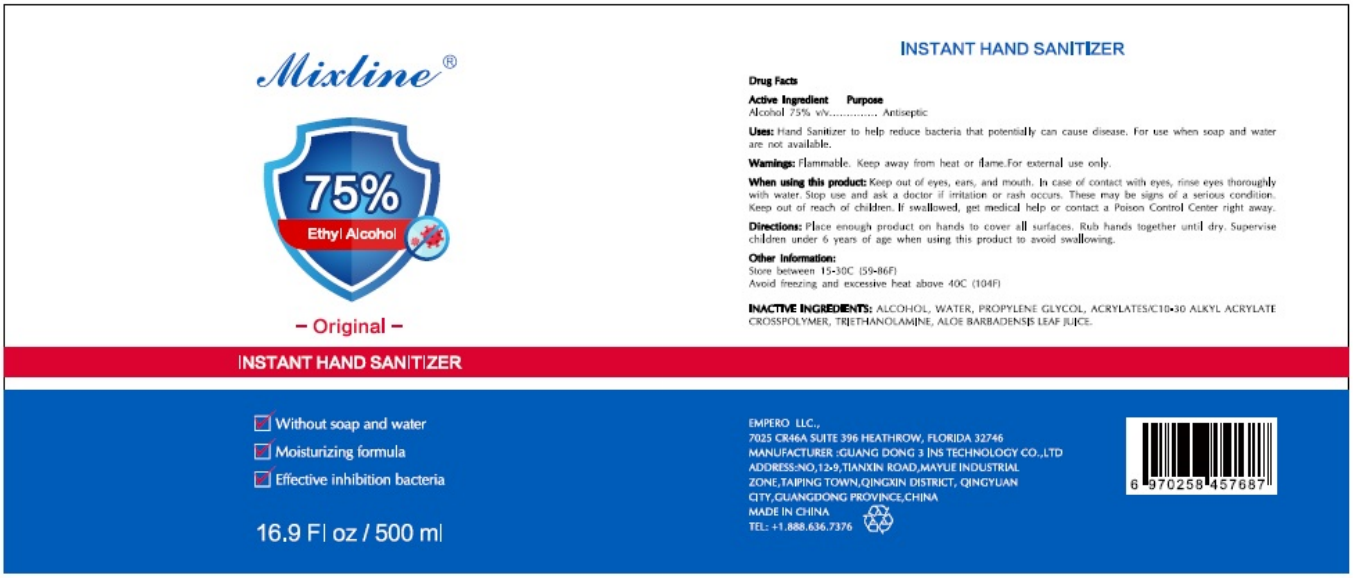
Mixline 75% Ethyl Alcohol

Original

INSTANT HAND SANITIZER

- Without soap and water
- Moisturizing formula
- Effective inhibition bacteria

16.9 Fl oz/ 500 ml



MIXLINE

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76778-075
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76778-075-17	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/28/2020	

Labeler - EMPERO LLC (024603167)

Registrant - EMPERO LLC (024603167)

Establishment

Name	Address	ID/FEI	Business Operations
GUANG DONG 3 INS TECHNOLOGY Co.,Ltd		552312961	manufacture(76778-075)

Revised: 4/2020 EMPERO LLC