

HAND SANITIZER- hand sanitizer gel
Ningbo Meiteli Cosmetic Co., Ltd.

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.

Hand Sanitizer

Active Ingredient(s)

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Use

Hand Sanitizer to help reduce bacteria on the skin.

Warning

Flammable. Keep away from heat or flame

For external use only.

Do not use near eyes

In case of contact rinse eyes thoroughly with water.

Stop use

Stop use and see a doctor if irritation or rash appears and lasts

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a poison control Center right away.

Directions

Place enough product in your palm and rub hands together briskly until dry
Children under 6 years of age should be supervised when using this product

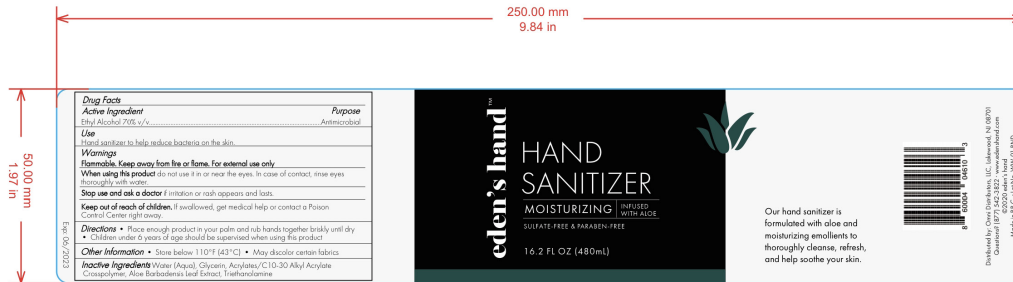
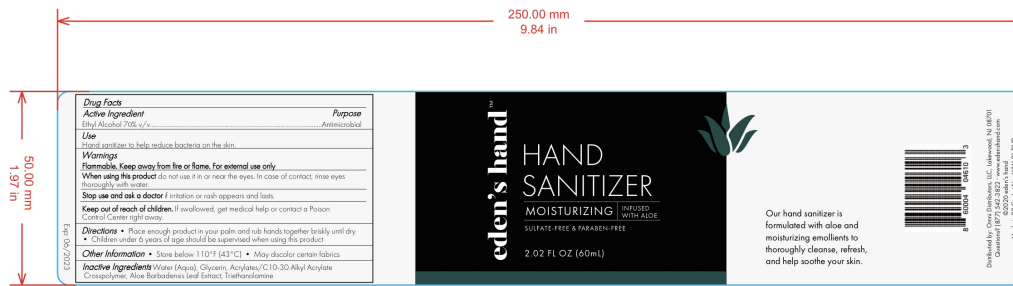
Other

- Store below 110F(43C)
- May discolor some fabrics or surfaces

Inactive ingredients

Water(Aqua), Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Extract, Triethanolamine

Package Label - Principal Display Panel



HAND SANITIZER

hand sanitizer gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:75541-007

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
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75541-007-03	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
2	NDC:75541-007-02	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	
3	NDC:75541-007-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/21/2020	

Labeler - Ningbo Meiteli Cosmetic Co., Ltd. (546624772)**Establishment**

Name	Address	ID/FEI	Business Operations
Ningbo Meiteli Cosmetic Co., Ltd.		546624772	manufacture(75541-007)

Revised: 6/2020

Ningbo Meiteli Cosmetic Co., Ltd.