

HUMASTE HAND SANITIZER- ethyl alcohol gel
Swanrose, 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.

HUMASTE Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antimicrobial

Uses

- Hand sanitizer to reduce microorganisms on the skin.
- Use this product when soap and water are not available.

Warnings

• **For external use only.** • Avoid contact with eyes. If contact occurs, rinse thoroughly with water. • **FLAMMABLE. This product contains ethyl alcohol. Keep away from sources of ignition.** • Discontinue use if irritation or redness develops. • If irritation persists for more than 72 hours, consult a physician.

• **KEEP OUT OF REACH OF CHILDREN.**

- If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- **Read the entire label before using this product.**
- Place enough product on your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, Glycerine, 1,2 Propylene Glycol, Carbomer, 2-Amino-2-Methyl Propanol

70% Alcohol
Self Drying Gel
Rub well until dry

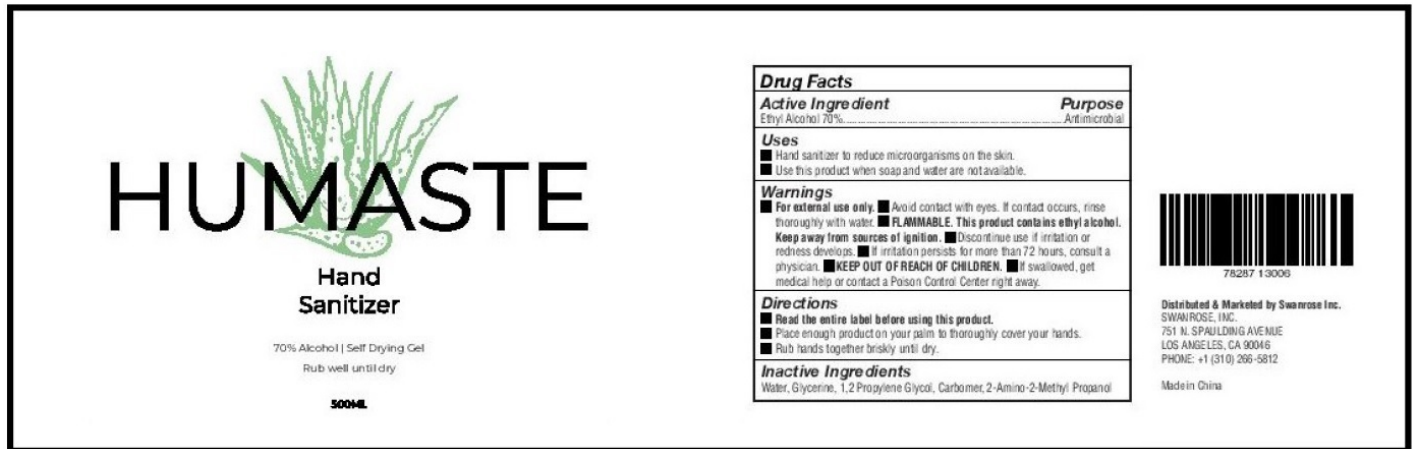
Distributed & Marketed by Swanrose Inc.

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Made in China

Packaging



HUMASTE HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78287-130
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HO MO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78287-130-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
2	NDC:78287-130-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
3	NDC:78287-130-03	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
4	NDC:78287-130-04	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
5	NDC:78287-130-05	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	

6	NDC:78287-130-06	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
7	NDC:78287-130-07	560 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
8	NDC:78287-130-08	980 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
9	NDC:78287-130-09	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
10	NDC:78287-130-10	4000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
11	NDC:78287-130-11	5000 mL in 1 CAN; Type 0: Not a Combination Product	06/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/25/2020	

Labeler - Swanrose, Inc. (117523391)

Establishment

Name	Address	ID/FEI	Business Operations
ZHEJIANG JINGHUI COSMETICS SHARE CO.,LTD		529558167	manufacture(78287-130)

Revised: 6/2020

Swanrose, Inc.