

VINIFERAMINE HAND SANITIZER- ethyl alcohol spray
MCCORD RESEARCH

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.

VINIFERAMINE HAND SANITIZER

Active Ingredient

Ethyl Alcohol

Purpose

Antiseptic

Uses

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

Warnings

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

Do not use

- in children less than 2 months of age
- on open skin wounds

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 15-30°C (59-86F)

Avoid freezing and excessive heat above 40°C (104F)

Inactive Ingredients

Water, Glycerin, Pyrus Malus (Apple) Fruit Extract, Fragrance

Product Label

NDC 71358-017-02

VINIFERAMINE®

Hand Sanitizer

Small Molecule Technology
Without Scientific Equal™



8 fl oz (237 mL)

Drug Facts

Active Ingredient	Purpose
Ethyl Alcohol 64% w/v	Antiseptic

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 (855) 312-8667 www.viniferamine.com
Order Number: 79836 R1662

VINIFERAMINE HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71358-017
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	640 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
APPLE (UNII: B423VGH5S9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71358-017-02	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/26/2020	

Labeler - MCCORD RESEARCH (010011284)

Establishment

Name	Address	ID/FEI	Business Operations
Wasatch Product Development, LLC		962452533	manufacture(71358-017)

Revised: 1/2022

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