

HAND SANITIZER- alcohol soap

NVE Pharmaceuticals

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.

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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.

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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

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel



**TEAR AT DOTTED LINE
TAMPER EVIDENT**
Do not use if seal is broken

Non-aerosol • Dye-free

KILLS 99.99% OF GERMS

*Kills 99.99% of Most Common Germs
That May Cause Illness*

**DO NOT
DRINK**



#8284 R.042320

Manufactured and Distributed by NVE Pharmaceuticals
Andover, NJ 07821 | Tel. 1-800-548-3546

Drug Facts (cont.)
Directions • Place enough product in your palm to thoroughly cover your hands • Rub hands together briskly until dry • Children under 6 years of age should be supervised when using this product
Other information • Store below 110°F (43°C) • May discolor certain fabrics or surfaces
Inactive ingredients Water (Aqua), Methylcellulose, Glycerin, Hydrogen Peroxide.
Questions or comments? call 1-800-548-3546 Monday through Friday 8:00 AM to 5:00 PM

Hand Sanitizer

**Alcohol Antiseptic
70% Topical Solution**

UNSCENTED

2 FL OZ (60 mL)

Drug Facts
Active ingredient Ethyl Alcohol 70% v/v.....
Purpose Antimicrobial
Uses • Hand sanitizer to help reduce bacteria on the skin that could cause disease • Recommended for repeated use
Warnings Flammable. Keep away from fire or flame.
For external use only
When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
Stop use and ask a doctor if irritation or rash appears and lasts
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. ▶

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60 mL NDC: 78832-729-02

HAND SANITIZER

alcohol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78832-729
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0K00R)	

Product Characteristics

Color		Score	
Shape		Size	

Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78832-729-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/13/2020	



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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333A	11/13/2020	

Labeler - NVE Pharmaceuticals (004129433)

Registrant - NVE Pharmaceuticals (004129433)

Establishment

Name	Address	ID/FEI	Business Operations
NVE Pharmaceuticals		004129433	manufacture(78832-729)