

NEUTRALIZER- alcohol gel

AWSM Marcin Szakoła

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 70% 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 6 years 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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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 12 years of age when using this product to avoid swallowing.

Other information

- Store between 10-26C
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

200 mL NDC: 79887-114-24



HAND SANITIZER GEL

70%

- ALCOHOL ANTISEPTIC 70%
- TOPICAL SOLUTION
- NON-STERILE SOLUTION

200ml

Active ingredient:
Alcohol 70% v/v

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

Warning:
For external use only. Flammable. Keep away from heat and flame.

Do not use:

- in children less than 6 months of age.
- on open skin wound.

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 for use:
Place enough product on hands to cover all surfaces. Rub hands together until dry. Supervise children under twelve years of age when using this product to avoid swallowing.

Other information:
Store between 10 and 26C
Avoid freezing and excessive heat above 40C

Inactive Ingredients: Alcohol Denat., Aqua, Acrylates Copolymer, Parfum, Glycerin, Panthenol, Aloe Barbadensis Leaf Extract, Triethanolamine, Sodium Benzoate, Potassium Sorb
Best before/ SN/ Expiration Date: On bottle

Manufactured for:
Wisconsin Wine
and Spirits INC



500 mL NDC: 79887-114-54



HAND SANITIZER GEL

70%

- ALCOHOL ANTISEPTIC 70%
- TOPICAL SOLUTION
- NON-STERILE SOLUTION

500ml

Active ingredient:
Alcohol 70% v/v

Use:
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Warning:
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Do not use:

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Best before/ SN/ Expiration Date: On bottle

Manufactured for:
Wisconsin Wine
and Spirits INC



alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79887-114
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
SODIUM BENZOATE (UNII: OJ245FE5EU)	
PANTHENOL (UNII: WW9CM0067Z)	
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
ALOE (UNII: V5VD430YW9)	1 mL in 100 mL
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79887-114-24	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/14/2020	
2	NDC:79887-114-54	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/14/2020	

Labeler - AWSM Marcin Szakoła (422446268)

Establishment

Name	Address	ID/FEI	Business Operations
AWSM Marcin Szakoła		422446268	manufacture(79887-114)

Revised: 4/2021

AWSM Marcin Szakoła