

NAMAHANDSANITIZER- alcohol solution
Cospro Development Corp.

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) (70%, 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
- c. distilled water

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 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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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, polysorbate 20, aloe barbadensis leaf juice, tocopheryl acetate, maltodextrin, purified water

Package Label - Principal Display Panel



nama hand sanitizer

.....
 eliminates 99.9%
 of most bacteria

 powered by nature
 infused with Aloe Vera,
 Vitamin E & Glycerin
 refreshing mint aroma

242 ml e 8.2 fl oz

Drug Facts	
Active ingredient Ethyl Alcohol 70%	Purpose Antiseptic
Use • Hand sanitizer to help reduce bacteria on the skin	
Warnings	
Flammable. Keep away from fire and flame	
For external use only	
When using this product do not use in or near eyes. If contact occurs, 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.	
Directions • Put enough product in your palm to cover hands and 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 Aqua (Deionized Water), Glycerin, Polysorbate 20, Parfum (Fragrance), Aloe Barbadosis Leaf Juice, Tocopheryl Acetate, Maltodextrin	
Questions? www.namanaturals.com	

nama Hand Sanitizer

Never be without personal protection!
 Shield your hands from germs naturally
 with our plant-based hand sanitizer.

Botanical extracts of aloe vera, vitamin E
 and also glycerin help keep your skin
 healthy and hydrated. The refreshing
 mint aroma helps to soothe and recharge.



Made in USA & Dist. by
Cospro Development Corp.
 Reading, PA 19601
namanaturals.com



alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46607-117
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
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0K00R)	
ALOE (UNII: V5VD430YW9)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46607-117-01	242 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/29/2020	
2	NDC:46607-117-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/29/2020	

Labeler - Cospro Development Corp. (785638821)

Registrant - Cospro Development Corp. (785638821)

Establishment

Name	Address	ID/FEI	Business Operations
Cospro Development Corp.		785638821	manufacture(46607-117) , pack(46607-117) , label(46607-117)

Revised: 4/2020

Cospro Development Corp.