HONEYBEEHANDSANITIZER- 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 rub manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Rub Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand rub 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. Sterile distilled water

Active Ingredient(s)

Ethyl Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer Spray

Use

Hand Sanitizer to help reduce bacteria on the skin.

Warnings

For external use only. Flammable. Keep away from fire and flame

Do not use

do not use in or near eyes. If contact occurs, rinse eyes thoroughly with water

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 this and all drugs 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 110F (43C)
- May discolor certain fabrics or surfaces

Inactive ingredients

deionized water, glycerin, polysorbate 20, aloe barbadensis leaf juice, citrus aurantium dulcis peel oil, tocopheryl acetate, maltodextrin, parfum

Package Label - Principal Display Panel





SANITIZER HAND

SPRAY

with aloe and vitamin

Kills 99.9% of Most Bacteria Refreshing Citrus Scent 70% Ethyl Alcohol

59 ml /2 fl oz.

Active ingredient

Use • Hand sanitizer to help reduce bac

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 last get medical help or contact a Poison Control Center right away Keep this and all drugs out of reach of children, if swallow

and rub hands together briskly until dry. • Children under 6 years Directions • Put enough product in your palm to cover hands

Other Information • Store Below 110°F (43°C)

Polysorbate 20, Aloe Barbadensis Leaf Juice, Citrus Aurantium nactive Ingredients Aqua (Deionized Water), Glycerin Dulcis (Orange) Peel Oil, Tocopheryl Acetate, Maltodextrin.

iestions or Comments? www.honeybeegardens.com

Never be without protection!

Shield your hands from germs naturally with our plant-based hand sanitizer. Infused with aloe protect your hands while leaving skin smooth vera, vitamin E and glycerin. Cleanse and and hydrated.







Dist. by:

Cospro Development Corp., Reading, PA 19601 www.honeybeegardens.com alcohol solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46607-120	
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			
MALTO DEXTRIN (UNII: 7CVR7L4A2D)				
ALOE (UNII: V5VD430 YW9)				
GLYCERIN (UNII: PDC6 A3C0 OX)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
WATER (UNII: 059QF0KO0R)				
ORANGE PEEL (UNII: TI9T76XD44)				

ı	P	ackaging			
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:46607-120-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/0 1/20 20	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	12/0 1/20 20		

Labeler - Cospro Development Corp (785638821)

Registrant - Cospro Development Corp (785638821)

Establishment				
Name	Address	ID/FEI	Business Operations	
Cospro Development Corp		785638821	manufacture(46607-120), pack(46607-120), label(46607-120)	

Revised: 12/2020 Cospro Development Corp