HAND SANITIZER- alcohol gel LibertyWare LLC

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.

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

for handwashing to decrease bacteria on the skin.

Recommended for repeated use

Warnings

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

Do not use

• in the eyes. In case of contact rinse eyes thoroughly with water.

In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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

- Apply products to hands
- Briskly rub hands together until dry
- Supervise children under 6 years in the use of this product

Other information

- Store below 100F (43C)
- May discolor certain fabrics

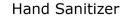
Inactive ingredients

distilled water, glycerin, rosemary, basil oil, aloe vera, vitamin e, azadirachtae indicae oil, essential oil,

Package Label - Principal Display Panel

250 mL NDC: 90076-100-01





Drug Facts

Active ingredients Ethyl Alcohol 70%

NDC Code: 90076-100-02

DUNS No.949700595

Uses • Hand Sanitizer to help to decrease bacteria on

Antiseptic

the skin Warnings · For external use only. Do not use in the eves, nose or mouth. In case of contact, rinse

thoroughly with water. Flammable. Keep away from fire or flame.

Stop use and ask a doctor if redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children • Children should only use under adult supervision

If swallowed seek medical attention or contact Poison

Directions • Dispense product into palms and thoroughly spread on the skin surfaces of both hands. Continue to rub until dry.

Other information • Store below 43°C (110°F). May discolor certain fabrics.

Inactive ingredients:

Distilled Water, Glycerine, Rosemary, Basil oil, Aloe Vera, Vitamin E, Azadirachtae Indiae Oil, Essential Oil, Carbomer 940, Tween 20.

Manufacturer: HCSP Company Limited No. 518, To Ngoc Van Street, Tam Phu Ward, Thu Duc District Ho Chi Minh City, Vietnam MADE IN VIETNAM

Distributed by: Libertyware LLC Clearfield, Utah, USA TEL: 888-500-5885











1000 mL NDC: 90076-100-02

NexSense Kills 99.9% of Germs Moisturizing with Aloe 34 fl. oz. (1000 ml)

Hand Sanitizer

Drug Facts

Active ingredients Ethyl Alcohol 70%

.Antiseptic

Uses • Hand Sanitizer to help to decrease bacteria on the skin

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DUNS No.949700595



Hand Sanitizer

Drug Facts

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Uses • Hand Sanitizer to help to decrease bacteria on the skin

Warnings • For external use only. Do not use in the eyes, nose or mouth. In case of contact, rinse thoroughly with water. Flammable. Keep away from fire or flame.

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NDC Code: 90076-100-02

DUNS No.949700595



HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:90076-100

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)
ALCOHOL (UNII: 3K9958 V90M) ALCOHOL 70 mL in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	1 mL in 100 mL			
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL			
ROSEMARY (UNII: IJ67X351P9)	1.45 mL in 100 mL			
WATER (UNII: 059QF0KO0R)	22.5 mL in 100 mL			
BASIL OIL (UNII: Z129 UMU8 LE)	1.45 mL in 100 mL			
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)	0.6 mL in 100 mL			
AZADIRACHTA INDICA SEED O IL (UNII: 4DKJ9B3K2T)	0.5 mL in 100 mL			
CARBOMER 940 (UNII: 4Q93RCW27E)	0.4 mL in 100 mL			
POLYSORBATE 20 (UNII: 7T1F30 V5YH)	0.1 mL in 100 mL			

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:90076-100-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			
2	NDC:90076-100- 02	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			
3	NDC:90076-100- 03	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	03/30/2020					

Labeler - LibertyWare LLC (949700595)

Registrant - LibertyWare LLC (949700595)

Establishment							
Name	Address	ID/FEI	Business Operations				
LibertyWare LLC		949700595	manufacture(90076-100)				

Revised: 10/2020 LibertyWare LLC