HAND SANITIZER- alcohol liquid Thomas Chemicals, 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.

Summers cape scented 70% hand sanitizer

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 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, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

59 mL NDC: 76865--032-01



SUMMERSCAPE

HAND SANITIZER

The crisp scent of fresh picked Strawberries from the vine mixed with a subtle blend of cucumber and honeydew.

Store in a cool dark area.

2 oz

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Purpose

Drug Facts

Active Ingredient[s]

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Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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.

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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: purified water, fragrance, glycerin, vitamin E, hydrogen peroxide



59 mL NDC:

SUMMERSCAPE

HAND SANITIZER LIQUID FORM 70%

The crisp scent of fresh picked Strawberries from the vine. Store in a cool dark area.



8 oz

Drug Facts
Active Ingredient[s]
Alcohol 70% v/v.
Use[s]
Health care personnel hand runb to help reduce bacteria that potentially can cause disease Warnings
For external use only. Flammable. Keep away from heat or flame. *Purpose*Antiseption

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Avoid Freezing and excessive heat above 40C (104F)
Inactive Ingredients purified water, fragrance, glycerin, vitamin E, hydrogen peroxide





76865-032-02

HAND SANITIZER

alcohol liquid

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Product	Information	

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76865-032

Route of Administration TOPICAL

ALPHA-TO COPHEROL ACETATE (UNII: 9E8X80D2L0)

Active Ingredient/Active Moiety

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Ingredient Name		Basis of Strength	Strength
I	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 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: 059QF0KOOR)

0.2 mL in 100 mL

l	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:76865-032- 01	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/18/2020	
	2 NDC:76865-032- 02	236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/18/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/18/2020	

Labeler - Thomas Chemicals, LLC (117504978)

Registrant - Jessica Tutt (117504978)

Establishment				
Name	Address	ID/FEI	Business Operations	
Thomas Chemicals, LLC		117504978	manufacture (76865-032)	

Revised: 8/2020 Thomas Chemicals, LLC