

JOURNEYMAN DISTILLERY HAND SANITIZER- ethanol solution

Journeyman Distillery, 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.

Journeyman Distillery Hand Sanitizer

Drug Facts

Active ingredient

Alcohol 80% v/v

Purpose

Antiseptic

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 immediately.

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 15-30°C (59-86°F)
- avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

JOURNEYMAN DISTILLERY

THREE OAKS, MI 49128

269-820-2050

JOURNEYMANDISTILLERY.COM

JOURNEYMAN Distillery

HAND SANITIZER

WARNING: NOT FOR CONSUMPTION

ALCOHOL ANTISEPTIC 80%

NON-STERILE TOPICAL SOLUTION

2 FL OZ / 59 ML

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77395-001
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
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77395-001-01	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/20/2020	

Labeler - Journeyman Distillery, LLC (013794196)

Establishment

Name	Address	ID/FEI	Business Operations
Journeyman Distillery, LLC		013794196	manufacture(77395-001)

Revised: 5/2020

Journeyman Distillery, LLC