

RED FOX HAND SANITIZER- red fox hand sanitizer liquid

Dehner 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.

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 80% v/v. Purpose: Antiseptic

HAND SANITIZER

• HAND SANITIZER • HAND SANITIZER



RED FOX

HAND SANITIZER

ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION

NON-STERILE SOLUTION

1.75L | 59.17FL OZ

PRODUCED & BOTTLED BY:

DEHNER DISTILLERY

CLIVE, IA 50325

DEHNERDISTILLERY.COM

Drug Facts

Active Ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

Use[s]

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.

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

~DO NOT CONSUME~

Manufactured in accordance
with WHO and FDA standards:
Emergency Supply.



HAND SANITIZER • HAND SANITIZER • HAND SANITIZER • HAND SANITIZER

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.

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

- 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

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	141.6 mL in 177 mL
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)			32.6 mL in 177 mL	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)			0.23 mL in 177 mL	
GLYCEROL FORMAL (UNII: 3L7GR2604E)			2.57 mL in 177 mL	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77305-0001-1	1750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2020	
2	NDC:77305-0001-2	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333A	05/05/2020	

Labeler - Dehner Distillery LLC (080131792)

Registrant - Joseph Dehner (080131792)

Establishment

Name	Address	ID/FEI	Business Operations
Dehner Distillery LLC		080131792	manufacture(77305-0001)

Revised: 5/2020

Dehner Distillery LLC